From: Sent: To: Subject:

Borin, Michael Friday, September 09, 2005 4:23 PM STIC-Biotech/ChemLib Search request: 10/617217

Examiner: M.Borin AU: 1631. Mailbox: 2C70 Office: Remsen 2A55 Tel.: 20713

RE: 10617217; EST

Please search nucleic acid SEQ ID NO: 88.

Thank you

ma 2717

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STAFF USE ONLY	Type of Search	Vendors and cost where applicable
Searcher: Searcher Phone: 2- Date Searcher Picked up: Date Completed: Searcher Prep/Rev. Time: Online Time:	NA#:AA#: Interference:SPDI: S/L:Oligomer: Encode/Transl: Structure#:Text: Inventor: Litigation:	STN: DIALOG: QUESTEL/ORBIT: LEXIS/NEXIS: SEQUENCE SYSTEM: LWWW/Internet: Other(Spp=5):
Date completed: Searcher: Bever Ly e Z	Search Site	Vendors IG
Terminal time:	CM-1	STN
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Total time:	N.A. Sequence	Geninfo
Number of Searches:	A.A. Sequence	SDC .
Number of Databases:	Structure	DARC/Questel
	Bibliographic	Other CGA



STIC Search Report Biotech-Chem Library

STIC Database Tracking Number 1985

TO: Michael Borin

Art Unit: 1631

Saaraa Natae

Location: REM/2A55/2C70

Serial Number: 10/617217

Friday, September 23, 2005

From: Beverly Shears

Location: Biotech-Chem Library

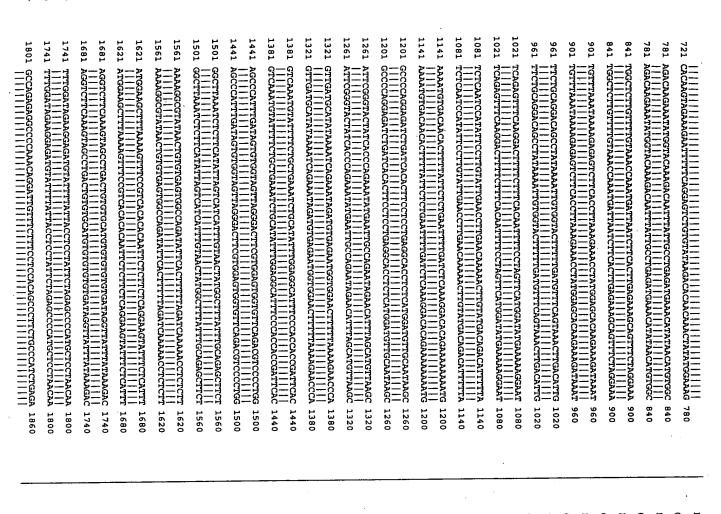
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Phone: 571-272-2528

beverly.shears@uspto.gov

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2340	2281 AAAATAAATTTTTATAGAGATCCTTGAGTAAAGACATTTTGCTTAATTTCTTTTTTCTTA	В
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RESULT 5
US-09-768-826-15
Sequence 15, Application 06/09768826
Fatent (0. US20020012966A1)
GENERAL MACHION
GENERAL SAI et al.
APPLICANT: Shi et al.
FITLE OF INVENTION: 18 human secreted proteins
FILE REFERENCE: PF512P1
CURRENT APPLICATION NUMBER: US/09/768,826
CURRENT FILING DATE: 2001-01-25

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; PRIOR APPLICATION NUMBER: PCT/US00/22350; PRIOR FILING DATE: 2000-08-15; PRIOR APPLICATION NUMBER: 60/148,759; PRIOR FILING DATE: 1999-08-16; NUMBER OF SEQ ID NOS: 61; SOFTWARE: Patentin Ver. 2.0; SEQ ID NO 15; LENGTH: 1860; TYPE: DNA ORGANISM: Homo sapiens
US-09-768-826-15
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Best Local Similarity 99.9%;
Matches 1837; Conservative
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Database: GenEmbl:* 1: gb ba:* 2: gb-htg:* 3: gb in:* 4: gb-om:* 5: gb-om:* 6: gb pat:* 7: gb-ph:* 8: gb pi:* 9: gb pr:* 10: gb ro:* 11: gb sts:* 12: gb-sy:* 14: gb_vi:*	Post-processing: Minimum Match 0% . Maximum Match 100% . Listing first 45 summaries	Minimum DB seq length: 0 Maximum DB seq length: 2000000000	Total number of hits satisfying chosen parameters: 9416466	Searched: 4708233 seqs, 24227607955 residues	Scoring table: IDENTITY_NUC Gapop 10.0 , Gapext 1.0	Title: US-10-617-217A-88 Perfect score: 2717 Sequence: 1 aaaaggaagacagaaaagccaataaatgtttttattcttt 271	Run on: September 22, 2005, 23:45:41 ; Search time 11587 Seconds (without alignments) 11362.115 Million cell updates/sec	OM nucleic - nucleic search, using sw model	Copyright (c) 1993 - 2005 Compugen Ltd.
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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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ALIGNMENTS

ORIGIN	PEATURES		AUTHORS TITLB JOURNAL COMMENT	VERSION KEYWORDS SOURCE ORGANISM	RESULT 1 BD170656 LOCUS DEFINITION ACCESSION
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/organism="nomo sapiens" /mol_type="genomic DNA" /db_xref="taxon:9606"	PC GOING CC NF-ke FH Key FT CDS	-AU 2N1 2N1 1K3 61P	Matsuda, A., Honda, G., Muramatsu, S. and Nagano, Y. NF-kappa B activation gene Patent: WO 02053737-A 44-11-5UL-2002; ASAHI KASEI CORP, AKIO MATSUDA, BOICHI HONDA, SHUJI MURAMATSU, YUKIKO NAGANO OS Homo sapiens (human) PN WO 02053737-A/44 PD 11-JUL-2002	BD170656.1 GI:27876468 WO 02053737-A/44. Homo sapiens (human) M Homo sapiens Eukaryota; Metazoa; Chordata; Cran Mammalia; Eutheria; Primates; Cata 1 (bases 1 to 2717)	BD170656 2717 bp DNA linear PAT 17-JAN-2003 N NF-kappa B activation gene.
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Wagano,Y., Doi,T., Shimotohno,K., Harada,T., N
and Sugaro,S.
Matsuda, A., Suzuki, Y., Honda, G., Muramatsu, S., Mutsuzaki, O., Nagano, Y., Harada, T., Nishida, E., Hayashi, H. and Sugano, S. Direct Submission
Submitted (25-NOV-2002) Yutaka Suzuki, The Institute of Medical Science, The University of Tokyo; Minatoku Shirokanedai 4-6-1, Tokyo, Tokyo 108-8639, Japan (E-mail:ysuzuki@ims.u-tokyo.ac.jp, Tel:81-3-5449-5343, Fax:81-3-5449-5416)
                                                                                                                                   Large-scale identification and characterization of activate NP-kappaB and MAPK signaling pathways Oncogene 22 (21), 3307-3318 (2003)
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Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Buteleostomi;
Mammalia Butheria; Primates; Catarrhini; Hominidae; Homo.
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1 (bases 1 to 3453)
Hardy,M.P., McGettrick,A.F., Ejdeback,M. and O'Neill,L.A.J.
Identification of a novel variant of the Toll-like receptor adaptor protein TIRAP3
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                            AATCTGCTACAAGATGACTTTGGTATCAAACCCGGAATAATCTTTGCTGAGATGCCATGT
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/gene="TIRAP3"
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/gene="TIRAP3"
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/protein_id="AAQ97432.1"
/protein_id="AAQ97432.1"
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/translation="MGIGKSKINSCPLS"/ILLIENFLADTWCMPGYTSLMS
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Consensus quality: 113349 bases at least Q30
Consensus quality: 113349 bases at least Q30
Consensus quality: 114811 bases at least Q30
Estimated insert size: 204000; pulse field gel estimation
Estimated insert size: 122110; sum-of-contigs estimation
Quality coverage: 7.16 in Q20 bases; pulse field gel estimation
Quality coverage: 11.95 in Q20 bases; sum-of-contigs estimation.
* NOTE: This is a 'working draft' sequence. It currently
* consists of 23 contigs. The true order of the pieces
* is not known and their order in this sequence is record is
* arbitrary Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
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Mammalia; Eutheria; Primates; Ca
1 (bases 1 to 124310)
DOE Joint Genome Institute.
Sequencing of Human Chromosome 5
Unpublished
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Submitted (03-AUG-1999) Production Sequencing Facility, DOE Joint
Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA
On Apr 20, 2001 this sequence version replaced gi:7708968.
------Genome Center
Center: Joint Genome Institute
Center: Ode: JGI
Web site: http://www.jgi.doe.gov
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Center Project Name:
Center clone name: Cl
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HTG; HTGS_PHASE1; HTGS_DRAFT; HTGS_ACTIVEFIN
Homo sapiens (human)
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/mol_type="genomic DNA"
/db_xref="taxon:9606"
/chromosome="5"
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21997: gap of unknown length
28247: contig of 6250 bp in length
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31864: contig of 3517 bp in length
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31964: gap of unknown length
35967: gap of unknown length
42256: contig of 6258 bp in length
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56435: contig of 7716 bp in length
64551: gap of unknown length
64551: contig of 7716 bp in length
64551: contig of 71807 bp in length
64551: contig of 1718 bp in length
64551: contig of 18016 bp in length
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                                                       TATTTGCAGAGCTTCTAAAAGGCGTATAACTGTGTGAGTGGCCA
                                                                                                                                                                                                CCCACCACCGATTCACAGCCCATTTGATAGTGTGGTAGTTAGGGACTTCC
                                                                                                                                                                                                                                                                                                      TITTIAAAAGAACCCAGICAAAIGIATITICIGCIGAAAICIGCATATITIGGAGGCATIT 1424
                                                                                                                                                                                                                                                                                                                                       GGATGTTGCAATAAGCATTCGGGTACTATCACCCAGAAATATGAATTGCCAGAATAGAAC
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                                                                      CTAATAAAACATGAACTGCCCACTCTTCATGCCTGCCAAACTTGGGGCAATTGATGCTAA
                                                                                                       CTAATAAAACATGAACTGCCCACTCTTCATGCCTGCCAAACTTGGGGGCAATTGATGCTAA 2684
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Submitted 128-FEB-2001) DOE Joint Genome Institute, 2800 Mitchell

Drive, Walnut Creek, CA 94598, USA

On Feb 28, 2001 this sequence version replaced gi:9256164.

Draft Sequence Produced by DOE Joint Genome Institute

www.jgi.doe.gov

rinishing Completed at Stanford Human Genome Center

www-shgc.stanford.edu

Quality: Phrap Quality >=40 99.6% of Sequence;

Estimated Total Number of Errors is 0.5.

STS Content:

WI-3489 G02883

SHGC-13356 G14683

SHGC-13164 G60385

WI-11806 G22104
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DOE Joint Genome Institute.
Direct Submission
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Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 147140)
DOB Joint Genome Institute and Stanford Human Genome Center.
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DOE Joint Genome Institute and Stanford Human Genome Center.
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 CAGGAGGGAGCTCAGAGCGTGGAAGAGATGTTTGAAGAAGAAGCTGAAGAAGAGGTGTTC
                     CAGGAGGGAGCTCAGAGCGTGGAAGAAGATGTTTGAAGAAGAAGCTGAAGAAGAAGATGTTTC
                                                                                           GATCTATCCTTGTGTAATGTTGCTGAGCACAGCAATACAACAGAGGGGCCAACAGGAAAG
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                                                                        GATCTATCCTTGTGTAATGTTGCTGAGCACAGCAATACAACAGAGGGGCCAACAGGAAAAG
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                                                                                                                                                                                                                                                                                                                                                                                                                                                         /mol_type="genomic_DNA"
/db_xref="taxon:9606"
/chromosome="5"
/clone="CTC-286N18"
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clone CTC-286N18, complete sequence.
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1245 GGATGTTGCAATAAGCATTCGGGTACTATCACCCAGAAATATGAATTGCCAGAATAGAAC 1304	1065 GATATGAAAAAGGAATTCTCAATCCATATTCCTTGTATTGAACCTTGAACAAAAACTTGT 1124	885 AGCAGTITCTAGGAAATGTTTAAATAAAAGAGAGTCTTCACCTTAAAGAAACCTATGGAG 944	705 AGTCGTGGATTTCCTACACAAGTAGAAGAATTTTTCAGGAGTCTGTGTATAAGACACAA 764	CCCCTTCCCCGAGAAAGGACTCCCTTTGCCCTCCAAACCATGAAGGCCTTAGAGGAAGAA	345 CTCAAATTTGTGATATTGCATGCAGAAGATGACACAGATGAAGCCCTCAGAGTCCAGAAT 404
	OY 235 ANTITITIANAGCITIANAGCITANAAGAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAA	93450 AATTTCTGCCACTGAGATGAATGTAACTGTGAGGCAAACTATTACCCTCTATGAGGATGTGAA 2025 GAAATGAAAGGGTTGAATTGATGATGTAACTGTTAAAGGCTTTTTGTCCTCTATGAGGATGTGAA	1845 TTCTGCCCTAACAAGCCAGAGAGCATCGTCCACAGAACAGATCATGAGATCAGGATCATGAGATCAGGATCAGATCAGATCAGATCAGATCAGATCAGAAACAGATCAGATCAGATCAGATCAGATCAGATCAGATCAGATCAGATCAGATCAGATCAGATCAGAAAACAGATCAGAAACAGATCAGAAACAGATCAGAAACAGATCAGAAACAGATCAGAAACAGAATCAAGAACATCAAGAACATCAAGAACATCAAGAACATCAAGAACATCAAGAACATCAAGAACATCAAGAACATCAGAAAAAACACCTCTCCGCAACAACAGAACATCAAGAACATCAAGAACAAAAACACCTCTTCCGAGAACATCAAGAACATCAAGAACATCAAGAACATCAAGAACATCATCAAGAACATCATCAAGAACATCATCAAGAACATCATCAAGAACATCATCAAGAACATCATCAAGAACATCATCAAGAACATCATCAAGAACATCATCAAGAACATCATCAAGAACATCATCAAGAACATCATCAAGAACATCATCAAGAACATCATCAAGAACATCATCAAGAACATCATCATCAAGAACATCATCAAGAACATCATCATCAAGAACATCATCATCAAGAACATCATCAAGAACATCATCATCAAGAACATCATCATCAAGAACATCATCATCAAGAACATCATCATCATCAAGAACATCATCAAGAACATCATCATCAAGAACATCATCATCAAGAACATCATCATCATCAAGAACATCATCAAGAACATCATCAAGAACATCATCATCATCAAGAACATCATCATCAAGAACATCATCAAGAACATCATCAAGAACATCATCAAGAACATCATCAAGAACATCATCAAGAACATCATCAAGAAACATCAAGAAACATCAAGAAACATCAAGAAACAAAAAAAA	93090 ATCARAAACCTCTCTTATGGAAGCTTTAAAAGTTTCCGTTATACAATTCCTCCAAGTCACCAATTCCTCCAGAAGTAGCTCAAAACCTCCTCTCCAGAAGTAGCTCAAAACCTCCTCCTATCCAGAAGTTACCTGACAAAGTTACCTCAAAACTAAAAGTAGCCTGACTGTGTGTG	Qy 1425 CCCACCACTCACAGCCCATTTGATAGTGTAGGTAGTTAGGGACTTCGTGGAGTGGTG 1484

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                                                                                                                                                                         Consensus quality: 151184 bases at least Q40
Consensus quality: 153297 bases at least Q20
Consensus quality: 153815 bases at least Q20
Estimated insert size: 159350; agarose-fp estimation
Estimated insert size: 154776; sum-of-contigs estimation
Quality coverage: 5.9 in Q20 bases; agarose-fp estimation
Quality coverage: 6.07 in Q20 bases; sum-of-contigs estimation.
* NOTE: This is a 'working draft' sequence. It currently
* consists of 6 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
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Mammalia; Eutheria; Primates; Catarrhini; Hominidae;
1 (Dases 1 to 15276)
DOE Joint Genome Institute.
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Center Project Name: 441299
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DOE Joint Genome Institute.
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HTG; HTGS_PHASE1; HTGS_DRAFT; HTGS_ACTIVEFIN.
Homo sapiens (human)
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Homo sapiens chromosome 5 clone RP11-90P14, WORKING
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7359: contig of 7359 bp in length
7369: gap of unknown length
17306: contig of 9847 bp in length
17406: gap of unknown length
37466: contig of 20060 bp in length
37566: gap of unknown length
57351: contig of 19785 bp in length
57351: gap of unknown length
91524: contig of 34073 bp in length
91524: gap of unknown length
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/mol_type="genomic DNA"
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/chromosome="5"
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GAAGTATTTCTCATTTAGGTCTTCAAAGTAGCCTGACTGTGTGCATGTGTGTG	TATTTGCAGAGCTTCTAAAAGGCGTATAACTGTGTGAGTGGCCAGATATTCACTTTTTAG 1604	CCCACCACCGATTCACAGCCCATTTGATAGTGTGGTAGTTAGGGACTTCGTGGAGTGGTG 1484	ATTTAGCATGTTAAGCGTTGATGCATATAAAATCAGAAATAGATGTGAGAATGGTGGAAC 1364	ACACAGAAAAAAAAATGGCCCCAGGAGATCTGATCACACTTCCTCCTGAGGCACCTCTCAT 1244	GATATGAAAAAGGAATTCTCAATCCATATTCCTTGTAATGAACCAAAAACTTGT 1124	CACANGANGANTANATTTCTTGCAGGACCATTTTCTTTCACANTTTTCCTAGTTCATG 1064 AGTANACTTGACATTGTCAGAGTTTCANGGACTTTTTCTTTCACANTTTTCCTAGTTCATG 1064 AGTANACTTGACATTGTCAGAGTTTCANGGACTTTTCTTTCACANTTTTCCTAGTTCATG 1064 AGTANACTTGACATTGTCAGAGTTTCAAGGACTTTTCTTTCACAATTTTCCTAGTTCATG 44543
SULT 7 304584 AY304584 AY304584 FINITION Homo sapiens TIRAP3b (TIRAP3) mRNA, complete cds. CESSION AY304584 CESSION AY304584 CESSION AY304584 AY30458 CESSI				TOTTTTTGCAGGTTACCTCTTCATAGCCATGAACCAAAACGTTCTATGAGGAGCATGCAAG	6 8 6 6 6 6 6 6 6 6 6 6 6 6 6 6 6 6 6 6	4 GAAATGAAAGGGTTGAATTGATGATGGATCTCTAAAGGCTTTTGTCCTCTATGAGGATGTGAA 5 AAACTAGGGACCACAAAAGGGAACAAGCAAAAAAGTTTGGATTTCGATAAAGTGATATGTA

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528 Db 1147 Qy 588 Db 1207 Qy 648 Db 1267 Qy 708 Db 1327 Qy 768 Db	348	Oy 228 288 00 00 00 00 00 00 00 00 00 00 00 00 0	B & B & B & B	G & B & B & B
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AB055258

Macaca fascicularis brain cDNA, clone:QflA-AB055258
AB055258

AB055258.1 GI:12698110
fis (full insert sequence).
Macaca fascicularis (crab-eating macaque)
Macaca fascicularis (brab-eating macaque)
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Macaca fascicularis (crab-eating macaque)
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Best Local Similarity
Matches 2498; Conserv
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R. Site1: DraIII (CACTGTGTG)

R. Site2: DraIII (CACTGTGTG)

R. Site2: DraIII (CACCATGTGTG)

R. Site2: DraIII (CACCATGTG)

Description: 1st strand cDNA was primed with an oligo(dT) primer [ATGTGGCCTTTTTTTTTTTTTTTT]; double-stranded cDNA was synthesized using specific 5' and 3' primers and amplified by PCR. The PCR product was digested with Sfil and size selection was performed to exclude fragments <1.5kb.The Sfil-digested PCR product was cloned into distinct DraIII sites of pMEJES-FL3. XhoI sites just outside the DraIII sites can be used to isolate the cDNA insert. Libraries were constructed by Sugano et al. (University of Tokyo, Institute of Medical Science). Custom primer used for sequencing

(5' end primer [CTTCTGCTCTANAAGCTGCG];

3' end primer [CGACCTGCAGCTCCAGCACA]).
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       Infectious Diseases, Division of Genetic Resources; 23-1, Toyama 1-chome, Shinjuku-ku, Tokyo 162-8640, Japan (B-mail:khashi@nih.go.jp, URL:http://www.nih.go.jp/yoken/genebank/, Tel:81-3-5285-1111(ex.2120), Fax:81-3-5285-1181)
Lab host: TOP10
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Osada, N., Hida, M., Kusuda, J., Tanuma, R., Iseki, K., Terao, K., Suzuki, Y., Sugano, S. and Hashimoto, K. Isolation of full-length cDNA clones from macaque
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Hashimoto, K., Osada, N.,
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Direct Submission
Submitted (02-FEB-2001) Katsuyuki Hashimoto, National Institute
Titute (102-FEB-2001) Toyama
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     AGATTCTTGCCCTCTTTCTCTCTCTTGGGGTAAAAGGCACAGTGTGGATACAAGTCCAGG
                                                                                                                                                               AAATTCCTGCCCTCTTTCTCTCTCTCTGGGGTAAAAGGCACAGTGTGGATACAAGTCCAGG
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                                                               ATATCATGAGTCAGATTCCAAGAAGTCTGAAGACCTATCCTTGTGTAATGTTGCTGAGCA
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/protein_id="BAB21882.1"
/db_xref="GI:12698111"
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/clome_Tib="macaque brain cDNA li
/dev_stage="adult"
355__720
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/mol_tyse="mRANA"
/db xrefs="taxon:9541"
/clone="Qf1A-10357"
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1334 AAATCAGAAATAGATGTGAGAATGGTGGAACTTTTTAAAAGAACCCAGTCAAATGTATTT 1393 	1274 CACCCAGAAATATGAATTGCCAGAATAGAACATTTAGCATGTTAAGCGTTGATGCATATA 1333	1214 TGATCACACTTCCTGAGGCACCTCTCATGGATGTTGCAATAAGCATTCGGGTACTAT 1273	ACTTTTATTCTCTGAATTTTGATCTCAAAGGACACAGAAAAAAAA	1094 TCCTTGTATTGAACCTTGAACAAAAACTTGTATGACAGACA	4 GACTITICTITCACAATITITCCTAGTTCATGGATATGAAAAAGGAATTCTCAATCCATAT 	974 CCTATAAAATTGTGGTACTTTTTGATGTTTCAGTAAACTTGACATTGTCAGAGTTTCAAG 1033 	914 GAGAGTCTTCACCTTAAAGAAACCTATGGAGCACAAGAAAGA	854 GTAAACCAAATGATTAATCTTCACTTGAGAAAGCAGTTTCTAGGAAATGTTTAAATAAA	794 GGTACAAAGACAATTTATTGCCTGAGATGAAACATATAACATGTGGCTGGC	734 AATTTTTCAGGAGTCTGTGTATAAGACAACAACAACTATATGGAAAGACAAGAAAAATAT 793 	674 CCTCCAAACCATCAATGCCTTAGAGGAAGAAAGTCGTGGATTTCCTACACAAGTAGAAAG 733	614 CTCTGTTATACCCATGCGGCCCCTGAACAATCCCCTTTCCCCGAGAAAGGACTCCCTTTGC 673	554 GTGTAATTTCCAGTTCTATACGTCCCTAATGAACTCCGTTAACAGGCAGCATAAATACAA 613 	494 TGTAAATGGGTCTGCATGGACAATCTTATTACTGACTGAAAACTTTTTTAAGAGATACTTG 553	434 CGGAATAATCTTTGCTGAGATGCCATGTGGCAGACAGCATTTACAGAATTTAGATGATGC 493 [374 TGACACAGATGAAGCCCTCAGAGTCCAGAATCTGCTACAAGATGACTTTGGTATCAAACC 433	314 GTTTGAAGAAGAAGAAGAAGAGGTGTTCCTCAAATTTGTGATATTGCATGCA	241 CAGCAATATAACAGAGGGGTCAACAAGAAAGCAGGAGGGAG
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Query Match Best Local S Matches 985 Qy 42 Db 146 Qy 102	CDS	PUBMED REFERENCE AUTHORS TITLE JOURNAL FEATURES	REFERENCE AUTHORS TITLE JOURNAL	- ACCESSION VERSION VERSION KEYWORDS SOURCE ORGANISM	Db RESULT 9 AY275836 LOCUS	5
y Match J6.1%; Score 981.4; DB 9; Length 1149; Local Similarity 99.4%; Pred. No. 9.4e-192; hes 985; Conservative 0; Mismatches 6; Indels 0; Gaps 0; 42 TCGCCTGCAGATTGAAAAGAAATGCTGAGAAATACATAAAAGTTTTCCTCTTCTGCCTTGG 101	//OL_Type="mrkh" //mol_Type="mrkh" /mol_Type="mrkh" /protein_id="mAP81748.1" /protein_id="makenglessessessessessessessessessessessessess	0 0		N Homo sapiens TIR domain-containing adapter protein mRNA, cds. AY275836 AY275836.1 GI:32435942 Homo sapiens (human) M Homo sapiens	ATT	2471 ACTTATTAGAGGAGATGGTTTTCATTGCATAGTGACATTTTCTTAGCCTTAACGTTC 2530
RESULT 10 CQ783319 LOCUS DEFINITION ACCESSION VERSION VERSION KEYWORDS SOURCE ORGANISM	B & B & B	\$ \$ \$ \$ \$	Qy Db	Q	Q	}
CQ783319 Sequence 3459 from Patent EP1396543. CQ783319 CQ783319 CQ783319 CQ783319 Homo sapiens (human) Homo sapiens Homo sapiens Bukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;	926 GAAACATTGACATTGACATTGACAATGACTAAAAACCTAAAACCTATAAACCTTCACTTTCACTTTCACTTTCACTTTCACTTTCACTTTCACTTTCACTTTCACTTTCACTTTCACTTTCACTTTCACTTTCACCTTTAAAAAA	702 GAAAGTCGTGGATTTCCTACACAAGTAGAAAGAATTTTTCAGGAGTCTGTGTATAAGACA 761		566 GGCAGACAGCATTTACAGAATTTAGATGCTGTAAAATGGGTCTGCATAGGACAATCTTA 625 522 TTACTGACTGAAAACTTTTTAAGAGATACTTTGGTGTAAATTTCCAGTTCTATACGTCCCTA 581 626 TTACTGACTGAAAACTTTTTAAGAGATACTTGGTGTAATTTCCAGTTCTATACGTCCCTA 685 626 TTACTGACTGAAAACTTTTAAGAGATACTTGGTGTAATTTCCAGTTCTATACCTCCCTA 685 682 ATGAACTCCGTTAACAGGCAGCATAAATACAACTCTGTTATACCCATGCGGCCCCTGAAC 641	402 AATCTGCTACAAGATGACTTTGGTATCAAACCCGGAATAATCTTTGCTGAGATGCCATGT 461 506 AATCTGCTACAAGATGACTTTGGTATCAAACCCGGAATAATCTTTGCTGAGATGCCATGT 461 506 AATCTGCTACAAGATGACTTTGGTATCAAACCCGGAATAATCTTTGCTGAGATGCCATGT 565 462 GGCAGACAGCATTTACAGAATTTAGATGATGATGTAAATGGGTCTGCATGGACAATCTTA 521	ATATTTATAATGGGTATCGGGAAGTCTAAAATAAATTCCTGCCCTCTTTCTCTCTC

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                                                                                                         ATGAACTCCGTTAACAGGCAGCATAAATACAACTCTGTTATACCCATGCGGCCCCTGAAC 641
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GAAAGTCGTGGATTTCCTACACAAGTAGAAAGAATTTTTCAGGAGTCTGTGTATAAGACA
                           AATCCCCTTCCCCGAGAAAGGACTCCCTTTGCCCTCCAAACCATCAATGCCTTAGAGGAA
                                          AATCCCCTTCCCCGAGAAAGGACTCCCTTTGCCCTCCAAACCATCAATGCCTTAGAGGAA 701
                                                                                        ATGAACTCCGTTAACAGGCAGCATAAATACAACTCTGTTATACCCATGCGGCCCCTGAAC
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nilarity 99.5%;
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/protein_id="CAP86531.1"
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NGTYKFCFSNBFSTFTIKTVYFDFQVGSDPPLFPSENRVSALTQMESACVSIHBALKS
VIDYQTHFRLREAQGRSRAEDLNTRVAY"
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/mol_type="unassigned DNA"
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0; Mismatches 4;
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PN JP 2002017375-A/3017
PD 22-JAN-2002
PF 07-JUL-2000 JP 2000253172
PI TOSHIO OTA, TETSUO NISHIKAWA, T
PI ISHII,
PI YURI KAWAI, AI WAKAWATSU, TOMOY,
SHINICHI KOJIWA,
PI TETSUJI OTSUKI, HISASHI KOGA
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/mol_type="genomic DNA"
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Pred. No. 5.2e-171;
0; Mismatches 4;
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GAAAGCAGTTTCTAGGAAATGTTTAAATAAAAGAGAGTCTTCACCTT
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PC C12P21/02,C12Q1/68//C12P21/08,G06F17/30,C12N15/00,C12N5/00 Primer for synthesizing full-length cDNA and use thereof FH k Location/Qualifiers FT CDS (245) (808) Primer for synthesizing full-length cDNA and Patent: JP 2002017375-A 3017 22-JAN-2002; HELIX RESEARCH INSTITUTE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Buteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

1 (bases 1 to 1687)

Ota,T., Nishikawa,T., Isogai,T., Hayashi,K., Ishii,S., Kawai,Y., Wakamatsu,A., Sugiyama,T., Nagai,K., Kojima,S., Otsuki,T. and C12N15/09, C07K14/47, C07K16/18, C12N1/15, C12N1/19, C12N1/21, C12N5/ YURI KAWAI, AI WAKAMATSU, TOMOYASU SUGIYAMA, KEIICHI NAGAI, 07-JUL-2000 JP 2000253172 TOSHIO OTA,TETSUO NISHIKAWA,TAKAO ISOGAI,KOJI HAYASHI,SHIZUKO 1687 bp DNA linear PAT 1 full-length cDNA and use thereof. use thereof PAT 18-SEP-2002 g ဂ္ဂ

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2 (bases 1 to 1687)
Isogai, T. and Otsuki, T.
Direct Submission
                                                                                 Isogai,T., Ota,T., Nishikawa,T., Hayashi,K., Otsuki,T.,
Suzuki,Y., Nagai,K., Sugano,S., Ishii,S.,
Kawai-Hio,Y., Saito,K., Yamamoto,J., Wakamatsu,A., Namura,Y.,
Kojima,S., Nagahari,K., Masuho,Y., Ono,T., Okano,K., Yoshikawa,Y.,
Aotsuka,S., Sasaki,N., Hattori,A., Okumura,K., Iwayanagi,T. and
Ninomiya,K.
                                                                                                                                                                                                                                                                                                                     AK075218 1687 bp mRNA linear F
Homo Bapiens cDNA FLJ90737 fis, clone PLACE1010827, w
to COP-COATED VESICLE MEMBRANE PROTEIN 924 PRECURSOR.
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Homo sapiens (human)
                                                                   NEDO human cDNA sequencing
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Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
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     ATGAACTCCGTTAACAGGCAGCATAAATACAACTCTGTTATACCCCATGCGGCCCCTGAAC
                      ATGAACTCCGTTAACAGGCAGCATAAATACAACTCTGTTATACCCCATGCGGCCCCTGAAC
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akqcryediaqgtkctlepquttgghtdvdcrledpdgkvlykkmkxqydsptptakk
ngtykercenspestfethkttyvpdpqvgedpplppsenrvsaltqmesacvsihealks
vidyqthfrlreaqgrsraedlutrvay"
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/protein_id="BAC11479.1"
/db_xref="GI:22761165"
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/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="ptACB1010827"
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Oshiumi,H., Shida,K., Matsumoto,M. and Seya,T.
Direct Submission
Submitted (22-APR-2003) Hiroyuki Oshiumi, Osaka University,
Submitted (22-APR-2003) Hiroyuki Oshiumi, Osaka University,
Department of Biology, Graduate School of Science; Machikaneyama
1-1, Toyonaka, Osaka 560-0043, Japan
[E-mail:oshiumi@bio.sci.osaka-u.ac.jp, Tel:81-6-6850-5432,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      Oshiumi,H., Sasai,M., Shida,K., Fujita,T., Matsumoto,M. and STICAM-2: a bridging adapter recruiting to Toll-like receptor TICAM-1 that induces interferon-beta
J. Biol. Chem. (2003) In press
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/codon_start=1
/product="TIR-containing adapter molecule-2"
/protein_id="BAC98399.1"
/protein_id="BAC98399.1"
/db_xref="GI:37360902"
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VAEHSNTTEGPTGKQEGAQSVEEMFEEEAEEEVFLKFVILHAEDDTDEALRVQNLLQD
VGIKPGIIFAEMFCGRQHLQNLDDAVNGSAWTILLITENFLRDTWCNFQFYTSIANS
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Oshiumi,H., Sasai,M., Shida,K., Pujita,T., Matsumoto,M. and StTIR-containing Adapter Molecule (TICAM)-2, a Bridging Adapter Recruiting to Toll-like Receptor 4 TICAM-1 That Induces Interferon-(beta)
J. Biol. Chem. 278 (50), 49751-49762 (2003)
                                                                                        Homo sapiens (human)
Homo sapiens
Eukaryota, Metazoa, Chordata, Craniata, Vertebrata, Euteleostomi,
Mammalia, Eutheria, Primates, Catarrhini, Hominidae, Homo.
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2 (bases 1 to 725)
Coshiumi, H., Shida, K., Matsumoto, M. and Seya, T.
Direct Submission
Submitted (28-AUG-2002) Hiroyuki Oshiumi, Osaka Medical Center for Cancer and Cardiovascular Diseases, Department of Immunology; 1-3-3
Nakamichi, Higashinari-ku, Osaka, Osaka 537-0025, Japan
(E-mail:oshiumi@bio.sci.osaka-u.ac.jp, Tel:81-6-6973-1209,
Fax:81-6-6973-1209)
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VNRQHKYNSVIJPMFELNUMPLPRERTPFALQTINALEEESRGFPTQVERIFQESVYKTQ
QTIWKETRNMVQRQFIA"
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/mol_type="mRNA"
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Pred. No. 6.7e-139;
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Fitzgerald, K.A., Rowe, I
Monks, B., Pitha, P. and
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      2 (bases 1 to 708)
Fitzgerald, K.A., Rowe, D., Barnes, B., (Monke, B., 1tha, P. and Golenbock, D.T.
TRIF and TRAM: TLR adapter molecules:
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

1 (Dases 1 to 708)

Pitzgerald, K.A., Rowe, D.C., Barnes, B.J., Caffrey, D.R., Visintin, A., Latz, E., Monks, B., Pitha, P.M. and Golenbock, D.T.

LATZ, E., Monks, B., Pitha, P.M. and Golenbock, D.T.

LPS-TLR4 Signaling to IRF-3/7 and NF-{kappa}B Involves the Toll

Adapters TRAM and TRIF

J.Exp. Med. 198 (7), 1043-1055 (2003)
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Unpublished
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                                                                                                                                                                                                                                                                              map="5q15-q32"
|. .708
                                                                                                                                                                                                                  gene="TRAM"
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                                                                                                                                                                                          note="TLR adapter molecule"
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Best Local Sin Matches 708; Query Match

Similarity

ilarity 100.0%; Conservative

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26.1%;

Score 708; DB 9; L Pred. No. 2.2e-135; Mismatches

Length 708;

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Gaps

0

601 GGATTTCCTACACAAGIAGAAAGAATTTTTCAGGAGTCTGIGTATAAAG 771 ATATGGAAAGAGACAAGAAATATTGCCTGA
711 GGATTTCCTACACAAGTAGAAAGAATTTTTCAGGAGTCTGTGTATAAGACACACAACAAACT
651 CCCCGAGAAAGGACTCCCTTTGCCCTCCAAACCATCAATGCCTTAGAGGAAGAAGTCGT
591 GTTAACAGGCAGCATAAATACAACTCTGTTATACCCATGCGGCCCCTGAACAATCCCCTT
531 GAAAACTTTTTAAGAGATACTTGGTGTAATTTCCAGTTCTATACGTCCCTAATGAACTCC
471 CATTACAGAATTTAGATGATGATGATGAATGGGTCTGCATGGACAATCTTATTACTGACT
411 CAAGATGACTTTGGTATCAAACCCCGGAATAATCTTTGCTGAGATGCCATGTGGCAGACAG
351 TTTGTGATATTGCATGCAGAAGATGACACAGATGAAGCCCTCAGAGTCCAGAATCTGCTA
291 GGAGCTCAGAGCGTGGAAGAGATGTTTGAAGAAGATGTGAAGAAGATGTTCCTCAAA
231 TCCTTGTGTAATGTTGCTGAGCACAGCAATACAACAGAGGGGCCAACAGGAAAGCAGGAG
171 CACAGTGTGGATACAAGTCCAGGATATCATGAGTCAGATTCCAAGAAGTCTGAAGATCTA
111 ATGGTTATCGGGAAGTCTAAAATAAATTCCTGCCCTCTTTCTCTCTC
ATGGGTATCO ATGGGTATCO ATGGGTATCO CACAGTGTGT CACAGTGTGT CACAGTGTGT TCCTTGTGTI GGAGCTCAGI TTTGTGATA TTTGTGATA CAAGATGAC CAAGATGAC CAAGATGAC

Search completed: Sept Job time : 11598 secs

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Result
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Maximum Match 100%
Listing first 45 summaries
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Maximum DB seq length: 200000000
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Perfect score:
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          Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.
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3: 95 c
6: 97 c
6: 97 c
7: 97 c
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Copyright (c) 1993 - 2005 Compugen Ltd.
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ALIGNMENTS

Query Match Best Local Sim Matches 874;	ORIGIN	FEATURES Bource	RESULT 1 BX391704/C LOCUS DEFINITION ACCESSION VERSION KEYWORDS SOURCE ORGANISM REFERENCE AUTHORS TITLE JOURNAL COMMENT
Query Match 31.0%; Score 841.2; DB 5; Length 904; Best Local Similarity 97.7%; Pred. No. 3.6e-185; Matches 874; Conservative 0; Mismatches 19; Indels 2; Gaps 2;	/organism="Homo sapiens" /mol_type="meNA" /mol_type="meNA" /db xref="taxon:9606" /clone="CS0D1030YG11" /tissue type="PLACENTA COT 25-NORMALIZED" /clone_lib="Homo sapiens PLACENTA COT 25-NORMALIZED" /note="Ist strand cDNA was primed with a NotI-oligo(dT) /primer. Five prime end enriched, double-strand cDNA was digested with Not I and cloned into the Not I and Ecor V sites of the pCMVSPORT 6 vector. Library was normalized."	Location/Qualifiers 1904	BX391704 Homo sapiens PLACENTA COT 25-NORMALIZED Homo sapiens cDNA clone CSODIOJOYGI1 3-PRIME, mRNA sequence. BX391704 Homo sapiens PLACENTA COT 25-NORMALIZED Homo sapiens cDNA clone CSODIOJOYGI1 3-PRIME, mRNA sequence. BX391704.2 GI:46875240 BX391704.2 GI:46875240

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RESULT 2
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                                                                                 BX397140
BX397140.2 GI:46846453
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
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Li,W.B., Gruber,C., Jessee,J. and Polayes,D.
Full-length cDNA libraries and normalization
Unpublished (2001)
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                                           TGTAACTGTGGGCAAACTATTTACCCTCCTTTATCTGTGAAATGAAAGGGTTGAATTGAT
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                                                                                 ACAAGCAAAAAGTGGGGAGGCGAGAAAGTGATAGGTAAGAGTGGCAGAAGGCGGGAGAT
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/clone Tib="Homo sapiens PLACENTA COT 25-NORMALIZED"
/note="Ist strand cDNA was primed with a NotI-oligo(dT)
/note="Ist strand cDNA was primed with a CDNA was
primer. Five prime end enriched, double-strand cDNA was
digested with Not I and cloned into the Not I and EcoR V
sites of the pCMVSPORT 6 vector. Library was normalized."
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           /organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
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                                                                                                                                                  National Cancer Institute / NIH
Bldg. 31 Rm10A07 Betheeda, MD 20892
Email: cgapbs-r@mail.nih.gov
Tissue procurement: Dr. Michael Brownstein
cDNA Library Preparation: Invitrogen Corp
cDNA Library Preparation: Invitrogen Corp
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Agencourt Bioscience Corporation
Clone distribution: MCC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: NDAM487 row: d column: 20
High quality sequence start: 26
High quality sequence stop: 629.
location/Qualifiers
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923 bp mRNA linear EST.

923 bp mRNA linear EST.
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NIH-MCC http://mgc.nci.nih.gov/.
NAtional Institutes of Health, Mammalian
Unpublished (1999)
Contact: Daniela S. Gerhard, Ph.D.
Office of Cancer Genomics
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      Eukaryota; Metazoa;
Mammalia; Eutheria;
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/organism="Homo sapiens"
/mol_rype="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:30399931"
/tissue_type="White Matter"
/dev_stage="Unknown"
/lab_host="BH10B-Ton A ( T1 a
/clone_lib="NIH_MGC_181"
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0; Mismatches 33;
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Best Local Similarity
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1 (bases 1 to 753)

Ebert, L., Heil, O., Hennig, S., Korn, B., Neubert, P., Partsch, E., Peters, M., Radelof, U. and Schneider, D.

I.M.A.G.E. CDNA Clone Collection
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   CR737757 Soares placenta Nb2HP Homo sapiens cDNA clone IMAGp971D037; IMAGE:132835 5', mRNA sequence.

CR737757 CR737757
CR737757.1 GI:51587215
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             REAPD Deutsches Ressourcenzentrum fuer Genomforschung
Heubnerweg 6, D-14059 Berlin, Germany
Tel: +49 30 32639 100
Fax: +49 30 32639 111
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       RZPD; IMAGD971D037.
RZPDLIB; I.M.A.G.E. cDNA Clone Collection;
Contact: Inge Arlart
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      This clone is available royalty-free from RZPD; contact RZPD (clone@rzpd.de) for further information. M13r, Primer sequence: TTCACACAGGAAACAGCTATGAC.

Location/Qualifiers
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Heubnerweg 6, D-14059 Berlin, Germany
Email: www.rzpd.de
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                RZPD Deutsches Ressourcenzentrum fuer Genomforschung
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Contact: Inge Arlart
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     Unpublished (2004)
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                                                                                                                                                                                                                                                          AGGAGATCTGATCACACTTCCTCCTGAGGCACCTCTCATGGATGTTGCAATAAGCATTCG
                                                                                                                                     TGCATATAAAATCAGAAATAGATGTGAGAATGGTGGAACTTTTTAAAAGAACCCAGTCAA 1385
       ATTTGATAGTGTGGTAGTTAGGGACTTCGTGGAGTGGTGTTCAGACGTCCCCTGGGGCCTT
                                                          ATGTATTTTCTGCTGAAATCTGCATATTTGGAGGCATTTCCCACCACCGATTCACAGCCC
                                                                                                                TGCATATAAAATCAGAAATAGATGTGAGAATGGTGGAACTTTTTAAAAGAACCCAGTCAA
                                                                                                                                                                                      GGTACTATCACCCAGAAATATGAATTGCCAGAATAGAACATTTAGCATGTTAAGCGTTGA
                                                                                                                                                                                                         GGTACTATCACCCAGAAATATGAATTGCCAGAATAGAACATTTAGCATGTTAAGCGTTGA
                                                                                                                                                                                                                                                                                                                                 Conservative
                                                                                                                                                                                                                                                                                                                                                                                                                    /organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGD971D037 ; IMAGE:132835"
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      /dev_stage="placenta obtained at birth (full term)
/lab_host="DH10B (ampicillin resistant)"
/clone_lib="Soares placenta Nb2HP"
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 'sex="Female"
                                                                                                                                                                                                                                                                                                                                                26.8%;
99.6%;
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                                                                                                                                                                                                                                                                                                                                 Score 727.4; DB 7;
Pred. No. 1.3e-158;
0; Mismatches 1;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         721
                                                                                                                                                                                                                      Email: cgapbs-r@mail.nih.gov
Tissue Procurement: Dr. Gary W. Hunninghake, U of I
Tissue Procurement: Dr. Gary W. Bento Soares, University of Iowa
CDNA Library Preparation: Dr. M. Bento Soares, University of Iowa
CDNA Sequencing by: Dr. M. Bento Soares, University of Iowa
DNA Sequencing by: Dr. M. Bento Soares, University of Iowa
Clone Distribution: Distribution information can be found at
Clone Distribution: Distribution/cgap.html
                                                                                                                                                                                                                                                                                                                                                                                  Bukaryota; Metazoa; Chordata; Craniata; Vertebrata; Buteleostomi; Mammalia; Butheria; Primates; Catarrhini; Hominidae; Homo.

1 (bases 1 to 728)

NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.

National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Homo sapiens
                                                                                                                                                                                                                                                                                                                                                      Contact: Robert
                                                                                                                                                                                                                                                                                                                                                                   Unpublished (1997)
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POLYA=Yes.
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/organism="Homo sapiens"
/mol_type="mcNA,"
/db_xref="taxon:9606"
/clone="UI-H-FT2-bjf-a-01-0-UI"
/tissue_type="Alveolar Macrophage"
/dev_stage="Adult"
/lab_host="DH10B (Life Technologies)"
/clone_lib="NCI_CGAP_FT2"
                                                                                                                                                                       ocation/Qualifiers
                                                                                                                                                                                                                                                                                                                                                      Strausberg, Ph.D.
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05-AUG-2004

1923

660

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420 1625 1565

300

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1685

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U of I es, University of Iowa

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Query Match
Best Local
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                                                                                           GAAGAAAGTCGTGGATTTCCTACACAAGTAGAAAGAATTTTTTCAGGAGTCTGTGTATAAG
                                                                                                                                                                                                                                                                                                                 CTAATGAACTCCGTTAACAGGCAGCATAAATACAACTCTGTTATACCCCATGCGGCCCCTG
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ACACAACAAACTATATGGAAAGAGACAAGAAATATGGTACAAAGACAATTTATTGCCTGA 818
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     Conservative
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Pred. No. 8.1e-156;
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Ebert, L., Heil, O., Hennig, S., N. Radelof, U., Schneider, D. and Ko: Human UnigeneSet - RZPD3
Unpublished (2003)
Contact: Ina Rolfs
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          879 TGAGAAAGCAGTTTCTAGGAAATGTTTAAATAAAAGAGAGTCTTCACCCTTAAAGAAA 935
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             This clone is available royalty-free from RZPD; contact RZPD (clone@rzpd.de) for further information. M13r, Primer sequence: TTTCACACAGGAAACAGCTATGAC.

Location/Qualifiers
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      Heubnerweg 6, D-14059 Berlin, Germany
Tel: +49 30 32639 101
Pax: +49 30 32639 111
www.rzpd.de
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               RZPD; IMAGP998E09130.

RZPDLIB; I.M.A.G.E. CDNA Clone Collection;
Human UniqueneSet - RZPD3 (RZPDLIB No.972)

http://www.rzpd.de/CloneCards/cgi-
bin/showLib.pl.cgi/response?libNo=972 Contact: Ina Rolfs
RZPD Deutsches Ressourcenzentrum fuer Genomforschung GmbH
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Im Neuenheimer Feld 580, D-69120 Heidelberg, Germany
 TATTTATAAAGACTTTGGATAGAAGGAGATGTATTTTATTACCTCCTATTCTAGAGCCCC
                         TATTTATAAAGACTTTGGATAGAAGGAGATGTATTTTATTACCTCCTATTCTAGAGCCCCC 1787
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                                                                            Conservative
                                                                                                                                                                                 /db xref="taxon:9606"
/clone="IMAGp998E09130 ; IMAGB:127472"
/sex="male"
                                                                                                                                                                   constructed by Bento Soares and M.Fatima Bonaldo."
                                                                                                                                                                                                                                                                                                                                                                                                                                                  /organism="Homo sapiens"
/mol_type="mRNA"
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                                                                                            26.0%;
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                                                                        Score 707.4; DB 5;
Pred. No. 5.9e-154;
0; Mismatches 2;
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                      Tunor Gene Index

Unpublished (1997)
Contact: Robert Strausberg, Ph.D.
Email: cyapbs-r@mail.nih.gov
Tissue Procurement: Dr. Jose Mercuende
CDNA Library preparation: Dr. M. Bento Soares, University of Iowa
cDNA Library Arrayed by: Dr. M. Bento Soares, University of Iowa
cDNA Library Arrayed by: Dr. M. Bento Soares, University of Iowa
cDNA Sequencing by: Dr. M. Bento Soares, University of Iowa
Clone Distribution: Clone distribution information can be obtained
from Dr. M. Bento Soares, bento-soares@uiowa.edu
The following repetitive elements were found in this cDNA
sequence: 1-47, AAT rich#Low_complexity (matched compliment)
Seq primer: Mil FORMARD
POLYA=Yes.
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UI-H-EDO-axc-b-08-0-UI.81 NCI CGAP_EDO Homo sapiens cDNA clone
UI-H-EDO-axc-b-08-0-UI 3', mRNA sequence.
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CA440665.1 GI:24805085
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1 (bases 1 to 681)

NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.

National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
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            TAATGCTTCCTGTAAAGCCAATTTTATATACTAATAAAACATGAACTGCCCACTCTTCAT
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/clone lib="NCI CGAP EDO"
/note="Organ: Left Pubic Bone; Vector: pT7T3-Pac
/note="Organ: Left Pubic Bone; Vector: pT7T3-Pac
/note="Organ: Left Pubic Bone; Vector: pT7T3-Pac
(Pharmacia) with a modified polylinker; Site_1: EcoR I;
Site_2: Not I; NCI CGAP_EDO is a cDNA library containing
the following tissue(s): Chondrosarcoma cell line CS5. The
library was constructed according to Bonaldo, Lennon and
Soares, Genome Research, 6:791-806, 1996. First strand
CDNA synthesis was primed with an oligo-dTT primer
containing a Not I site. Double stranded cDNA was ligated
to an EcoR I adaptor, digested with Not I, and cloned
directionally into pT7T3-Pac vector. The oligonucleotide
used to prime the synthesis of first-strand cDNA contains
a library tag sequence that is located between the Not I
site and the (dT)18 tail. The sequence tag for this
library is GCTCAAGGCT.
TAG_INSUB=chondrosarcoma
TAG_LIB=UI-H-EDO
TAG_SEQ=CGTCAAGGCT"
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/db_xref="taxon:9606"
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/lab_host="DH10B (Life Technologies)"
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Pred. No. 1.4e-143;
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Best Local
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                                                                                      cch 24.4%; al Similarity 99.1%; 676; Conservative
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          cDNA Library preparation: Dr. M. Bento Soares, University of Iowa cDNA Library Arrayed by: Dr. M. Bento Soares, University of Iowa DNA Sequencing by: Dr. M. Bento Soares, University of Iowa Clone Distribution: Clone distribution information can be obtained from Dr. M. Bento Soares, bento-soares@uiowa.edu

The following repetitive elements were found in this cDNA sequence: 1-47, >AT_rich#Low_complexity (matched compliment)
Seq primer: M13 FORWARD
POLYA=Yes.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    CA438249
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UI-H-DT1-avw-e-13-0-UI.s1 NCI_CGAP_DT1 Homo
UI-H-DT1-avw-e-13-0-UI 3', mRNA sequence.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Contact: Robert Strausberg, Ph.D.
Email: cgapbs-r@mail.nih.gov
Tissue Procurement: Dr. Jose Mercuende
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Tumor Gene Index
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National Cancer Institute, Cancer Genome Anat
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
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          GGTTGAATGATGGATCTCTAAAGGC-TTTGTCCTCTATGAGGATGTGAAAAAACTAGGGAC
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                                                                                                                                                                                                                                                            /clone lib="NCI CGAP DT1"

/clone lib="NCI CGAP DT1"

/note="Organ: Lung; Vector: pT7T3-Pac (Pharmacia) with a modified polylinker; Site_1: EcoR I; Site_2: Not I; NCI CGAP DT1 is a normalized cDNA library-containing the following tissue(s): Metatastic Chondrosarcoma in Lung. The library was constructed according to Bonaldo, Lennon and Soares, Genome Research, 6:791-806, 1996. First strand cDNA synthesis was primed with an oligo-dT primer containing a Not I site. Double stranded cDNA was ligated to an EcoR I adaptor, digested with Not I, and cloned directionally into pT7T3-Pac vector. The oligonucleotide used to prime the synthesis of first-strand cDNA contains a library tag sequence that is located between the Not I site and the (dT).8 tail. The sequence tag for this
                                                                                                                                                                                                             library is AACTGTTCGG.
TAG_TISSUE=lung metatastic chondrosarcoma
TAG_LIB=UI-H-DT1
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/tissue_type="Metastatic Chondro
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/lab_host="DH10B (Life Technologies)"
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|mol_type="mRNA"
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                                                                                    Score 662; DB 6; 1
Pred. No. 2.3e-143;
0; Mismatches 5;
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Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

B 1 (bases 1 to 887)

B 1 (bases 1 to 887)

National Institutes of Health, Mammalian Gene Collection (MGC)

Institutes of Health, Mammalian Gene Collection (MGC)

Lipublished (1999)

Contact: Robert Strausberg, Ph.D.

Email: cgapbs-r@mail.nih.gov

Tissue Procurement: ATCC

CDNA Library Preparation: Life Technologies, Inc.

CDNA Library Arrayed by: The I.M.A.G.B. Consortium (LINL)

DNA Sequencing by: Agencourt Bioscience Corporation

Clone distribution: MGC clone distribution information can be found through the I.M.A.G.B. Consortium/LLNL at:

http://image.llnl.gov

Plate: LLAM13218 row: d column: 04

High quality sequence stop: 664.
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                                                                                                                                                                                 ATGAACTCCGTTAACAGGCAGCATANATACAACTCTGTTATACCCCATGCGGCCCCTGAAC
                                                                                                                                                                                                                                                                                                      GGCAGACAGCATTTACAGAATTTAGATGATGCTGTAAATGGGTCTGCATGGACAATCTTA
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                                                                           AAGAAAGTCGTGGGATTTCCTACACAAGTAGAAAGNATTTTTTTCAGGAGTCTGTGGTAT
                                                                                        AAGAAAGTCGT-GGATTTCCTACACAAGTAGAAAGAATTTTTCAGGAGTCTGTGTATAAG
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                                                                                                                                            AATCCCCTTCCCCGAGAAAGGACT-CCCTTTGCCCTCCAAACCATCAATG-CCTTAGAGG
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/lab_host="HH108 (phage-resistant)"
/clone_lib="NIH_MGC_70"
/note="Organ: pancreas; Vector: pCMV-SPORT6; Site_1: NotI;
Site_2: Sall; Cloned unidirectionally. Primer: Oligo dT.
Average insert size 1.1 kb. Library constructed by Life Technologies."
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/clone="IMAGE:6018003"
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Pred. No. 4.1e-140;
n: Mismatches 32;
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LOCUS
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515
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ATTACACTTAAAGTCACAGGAAAAAATATACATGCTTACTCAGGCTTTCTTAAAAATAA
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RESULT 10

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Query Match
Best Local Similarity
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UI-H-ED0-axl-a-05-0-UI.81 NCI_CGAP_ED0 Homo sapiens cDNA clone
UI-H-ED0-axl-a-05-0-UI 3', mRNA sequence.
CA441688
CA441688.1 GI:24806108
EST.
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The following repetitive elements were found in this cDNA sequence: 1-54, >AT rich#Low_complexity (matched compliment) seq primer: M13 FORWARD
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NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
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Tissue Procurement: Dr. Jose Mercuende
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       Tumor Gene Index
Unpublished (1997)
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ATTACACTTAAAGTCACAGGAAAAAAATATACATGCTTACTCAGGCTTTCTTAAAAAATAA 2287
                                                                                                                      Conservative (
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 /cloime lib=NCI GAP_BDO"
/cloime lib=NCI GAP_BDO"
/note="Organ: Left Pubic Bone; Vector: pT7T3-Pac
/note="Organ: Left Pubic Bone; Vector: pT7T3-Pac
/note="Organ: Left Pubic Bone; Vector: pT7T3-Pac
/pharmacia) with a modified polylinker; Site 1: BcoR I;
Site 2: Not I; NCI_CGAP_BDO is a cDNA library containing
the following tissue(s): Chondrosarcoma cell line CS5. The
library was constructed according to Bonaldo, Lemnon and
Soares, Genome Research, 6:791-806, 1996. First strand
CDNA synthesis was primed with an oligo-dT primer
containing a Not I site. Double stranded cDNA was ligated
to an EcoR I adaptor, digested with Not I, and clomed
directionally into pT7T3-Pac vector. The oligonucleotide
used to prime the synthesis of first-strand cDNA contains
a library tag sequence that is located between the Not I
site and the (dT)18 tail. The sequence tag for this
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    library is GCTCAAGGCT.
TAG TISSUE=chondrosarcoma
TAG_LIB=UI-H-ED0
TAG_SEQ=CGTCAAGGCT"
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/db_xref="taxon:9606"
/clone="UI-H-EDO-ax1-a-05-0-UI"
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/dev_stage="Adult"
/lab_host="DH10B (Life Technologies)"
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                                                                                                                                                                                                                Contact: Morton, C. C.
Departments of Pathology and Obstetrics,
Reproductive Biology
Brigham and Women's Hospital
75 Francis Street, Harvard Medical School
                                                                                 DNA sequencing and analyses were performed by National Institute of Health Intramural Sequencing Center (NISC; see http://www.nisc.nih.gov).

This clone is available royalty-free through LLNL; contact the IMAGE Consortium (info@image.llnl.gov) for further information. Plate: LLNM6328 row: N column: 11
                                                                                                                                                                                                                                                                                                                         Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalla; Eutheria; Primates; Catarrhini; Hominidae; Homo.

1 (bases 1 to 615)

Robertson,N.G., Khetarpal,U., Gutierrez-Espeleta,G.A., Bieber,F.R. and Morton,C.C.

Isolation of novel and known genes from a human fetal cochlear cDNA library using subtractive hybridization and differential screening Genomics 23, 42-50 (1994)
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 615 bp df99306.yl Morton Fetal Cochlea IMAGE:2540746 5', mRNA sequence.
                                                                                                                                                                                        Tel: 617 732 7980 Fax: 617 738 6996
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                                                                                                                                                                           Email: ccmorton@bics.bwh.harvard.edu
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                                                        primer: M13RP1 reverse primer
Location/Qualifiers
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
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                                                                                                   RESULT 12
BI493311/c
LOCUS
                           VERSION
KEYWORDS
 SOURCE
ORGANISM
                                                          ACCESSION
                                                                                     DEFINITION
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BI493311 BI493311.1 EST.

Homo sapiens (human)

BI493311 613 bp mFdf99g06.w1 Morton Fetal Cochlea Homo IMAGE:2540746 3', mRNA sequence.

RNA linear EST 28-AUG-2001 sapiens cDNA clone

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This clone is available royalty-free through LLNL; contact the IMAGE Consortium (info@mage.llnl.gov) for further information. Plate: LLAM6328 row: N column: 11
Seg primer: T7 primer.
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Brigham and Women's Hospital
75 Francis Street, Harvard Medical School,
Tel: 617 732 7980
Fax: 617 738 6996
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1 (bases 1 to 613)

Robertson, N.G., Khetarpal, U., Gutierrez-Espeleta, G.A., Bieber, F.R. and Morton, C.C.

Isolation of novel and known genes from a human fetal cochlear cDNA library using subtractive hybridization and differential screening Genomics 23, 42-50 (1994)
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 CTATTCTGTTAGTACTTATTAGAGGAGGAGATGGTTTTCATTGCATAGTGACATTTTCTT 2517
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/db_xref="taxon:9606"
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Unpublished (1999)
Contact: Daniela S. Gerhard, Ph.D.
Office of Cancer Genomics
National Cancer Institute / NIH
Bldg: 31 Rm10A07 Bethesda, MD 2089
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1 (bases 1 to 585)

NIH-MGC http://mgc.nci.nih.gov/.
National Institutes of Health, Mammalian Gene Collection (MGC)
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AGENCOURT 14535036 NIH MGC 191 Homo sapiens cDNA clone
IMAGE:30417403 5', mRNA sequence.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      Email: cgapbs-r@mail.nih.gov
Tissue Procurement: Narayan Bhat
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http://image.llnl.gov
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            /organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/db_xref="taxon:9606"
/clone="IMAGE:30417403"
/tissue_type="pooled"
/lab_nost="PHIOB (TI phage-resistant)"
/lab_nost="PHIOB (TI phage-resistant)"
/clone_lib="NIH MGC_191"
/site_2: SfiI (ggccgctcggcc); Library is oligo-dT primed and directionally cloned. PBMC - Peripheral Blood Mononuclear Cells. RNA was pooled from 3/6hour stimulation with pMA adm Innomycin. 5' adaptor sequence:
5'-CACGGCCATTATGGCC-3' and 3' adaptors were used in cloning as follows: 5' adaptor sequence:
5'-CACGGCCATTATGGCC-3' and 3' adaptors sequence:
5'-CACGGCCATGAGGGCCGAACATG-dT(30)BN-3' (where B = A, C, or G and N = A, C, G, or T). Average insert size 1.69 kb (range 0.70-5.0 kb). 15/15 colonies contained inserts by PCR. This library was enriched for full-length clones and was constructed by Clontech Laboratories (Palo Alto, CA). Note: this is a NIH_MGC Library."
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Email: hbe@tigr.org
Clones are derived from the human BAC library RPCI-11.
library availability, please contact Pieter de Jong
                                                                                      Contact: Shaying Zhao, William Nierman, Department of Eukaryotic Genomics The Institute for Genomic Research 9712 Medical Center Dr., Rockville, MD 7e1: 301 838 0200 Fax: 301 838 0208
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                                                                                                                                                                                                                                                                                                                                                          Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; 1 (bases 1 to 560)
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/mol type="genomic DNA"
/db_xref="GDB:7541995"
/db_xref="taxon:9606"
/clone="RPCI-11-110F20"
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/clone_lib="RPCI-11"
/clone_totor: pBAce3.6; Site_1: EcoRI; Site_2: EcoRI;
/note="Vector: pBAce3.6; Site_1"
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Qy 2544 Db 173	2 4	Qy 2424 Db 293	Qy 2364 Db 353	Qy 2304 Db 413	Qy 2244 Db 473	Qy 2184 Db 533	Query Match Best Local Si Matches 533;	ORIGIN	FEATURES		JOURNAL UI	REFERENCE 1 AUTHORS NOTITLE NOTITLE	ORGANISM Homo
CTACTCACTTCTCTTTTCAGTTTTCATAATAAGTATTCATTTTTTTGCCATAATGCTTC 2603	GGAGATGGTTTTCATTGCATAGTGACATTTTCTTAGCCTTAACGTTCTGATAGTAGCTTA 2543 	GTTCTATGAGGAGCATGCAAGTAAGTCAAGCCTCCTATTCTGTTAGTACTTATTAGAGGA 2483 	CAGTACCGGGATCTGCACACATCTTTTTGCAGTTACCTCTTCATAGCCATGAACCAAAAC 2423 	TIGAGIAAAGACATITIGCTIAAITICTITITICTIAITCCCCACTIGIAIATCCCCTAC 2363 	CAGGAAAAAATATACATGCTTACTCAGGCTTTCTTAAAAATAAAT	ATATTTTTGTATATTGACAGCATAATTTATTTTTAATGCTGTCATTACACTTAAAGTCA 2243 	19.6%; Score 533; DB 2; Length 533; Similarity 100.0%; Pred. No. 2.8e-113; 3; Conservative 0; Mismatches 0; Indels 0; Gaps 0;		eq primer: -400P from Gibco igh quality sequence stop: 487. Location/Qualifiers 1	Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R. Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R. Emmert-Buck, M.D., Ph.D. CDNA Library Preparation: M. Bento Soares, Ph.D. CDNA Library Arrayed by: Greg Lennon, Ph.D. DNA Sequencing by: Washington University Genome Sequencing Center Clone distribution: NCI-CCAP clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL, send email to: info@image.llnl.gov	umbt gele index npublished (1997) ontact: Robert Strausberg, Ph.D.	<pre>ukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; ammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo. (bases 1 to 533) CI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap. ational Cancer Institute, Cancer Genome Anatomy Project (CGAP),</pre>	Homo sapiens

Search completed: September 23, 2005, 08:29:10 Job time : 8591 secs

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SUMMARIES

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NF-approximatelykB activating gene and expressed protein, applicable diagnosis and screening inhibitors or promoters to control excessive activation or inhibition for treating e.g. inflammations, autoimmune diseases and cancer.

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The invention relates to a purified protein (I), comprising one of 90

Claim 4; Page 403-407; 841pp; Japanese

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1.7	1.7	1.7	1.7	1.7	1.7	1.7	1.7	1.7	1.7	1.7	1.7	1.7	1.8	1.8	1.8	1.8	1.8	1.8	1.8	1.8	1.8	1.8	1.8	
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ALIGNMENTS

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28-DEC-2000; 2000JP-00402288.
26-MAR-2001; 2001JP-00088912.
24-AUG-2001; 2001JP-00254018.
                                                                                                                                                                                                                                                                                            Human; NF-kB; nuclear factor kappa B; mouse; antiinflammatory; immunomodulator; cytostatic; antiinfective; osteopathic; nootropic; neuroprotective; anti-HIV; autoimmune disease; cancer; infection; bone disease; AIDS; neurodegenerative disease; ischaemic disorder; gene;
                                                                                         WPI; 2002-583617/62.
P-PSDB; ABP61467.
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           ATAGTTGCAGAAGGCTTTATATATGCTTATAATGAAAAG
                                             AMACTAGGGACCACAAAAGGGAACAAGCAAAAAAGTTTGGATTCGATAAAGTGATATGTA
                                                           AAACTAGGGACCACAAAAGGGAACAAGCAAAAAAGTTTGGATTACGATAAAGTGATATGTA
                                                                                            GAAATGAAAGGGTTGAATTGATGGATCTCTAAAGGCTTTTGTCCTCTATGAGGATGTGAA
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WO2003029271-A2

10-APR-2003

24-SEP-2002; 2002WO-US030474.

24-SEP-2001; 2001US-0324631P

(HYSE-) HYSEQ INC

Tang TY, Zhang J, Zhou P, Ghosh M, Haley-Vicente D, D Wang D, Ma Y, Drmanac RT; r, Zhao Asundi ۷ę, Wang J, Wang Z, Wehrman Weng G;

New polynucleotide and polypeptide useful for diagnosing, preventing or treating conditions such as neurodegenerative diseases, anemias, platelet disorders, wounds, burns, ulcers, osteoporosis, autoimmune diseases or 2003-371981/35. DB; ADC31524.

Claim <u>.</u> SEQ ID ö 635; 1185pp; English.

The invention relates to 971 novel human cDNA sequences (ADC29919-ADC39889) and the polypeptides they encode (ADC30890-ADC31860). The invention also relates to nucleic acid sequences over 99% identical with the novel human cDNAs. The invention additionally encompasses expression

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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            cc recombinant production of a polypeptide of the invention; the cc against a polypeptide of the invention; a method of detecting cc against a polypeptide of the invention; a method of detecting cc polynuclectides or polypeptides of the invention; and methods of ci dentifying a compound which binds to a polypeptide of the invention. The cc invention further discloses methods of peventing, treating or ameliorating a medical condition, kits comprising polynuclectide probes condoration; methods for the identification of compounds that modulate the ci nevention; methods for the identification of compounds that modulate the contribution; which could be conting sequences corresponding to the CDNA sequences of the invention or cactivity of the polynucleotide and/or polypeptide; and 767 cc (ancist cation of mutations) properties of the invention are useful in diagnostics, drug screening, forensics, gene mapping, in the capacity of the coulcide acids and polypeptides of the invention are ct raits, for assessing biodiversity, and in producing many other types of data and products dependent on DNA and amino acid sequences. They are cf disease and other neurodegenerative diseases, anaemia, platelet cales used for treating diseases such as Parkinson's disease, Alzheimer's cf disease and other neurodegenerative diseases, anaemia, platelet cancer. The nucleic acids may also be used as hybridisation probes or concer. The nucleic acids may also be used as hybridisation probes or concer. The nucleic acids may also be used as hybridisation probes or concer. The nucleic acids may also be used as hybridisation probes or concer. The nucleic acids may also be used as hybridisation probes or concer. The nucleic acids may also be used as hybridisation probes or concer. The nucleic acids may also be used as hybridisation probes or concer. The nucleic acids may also be used as hybridisation probes or concer. The nucleic acids may also be used as hybridisation probes or concer. The polypeptides are also useful and the recombinant producti
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ACAACTCTGTTATACCCATGCGGCCCCTGAACAATCCCCTTCCCCGAGAAAGGACTCCCCT
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                                                                   CTTGGTGTAATTTCCAGTTCTATACGTCCCTAATGAACTCCGTTAACAGGCAGCATAAAT
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                                                                                                                                   CTATCACCCAGAAATATGAATTGCCAGAATAGAACATTTAGCATGTTAAGCGTTGATGCA
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XX PD O5-5 Human full-length cDNA, AAK94557 standard; 05-SEP-2001 EP1130094-A2 Homo sapiens Human; full length cDNA; cDNA synthesis; oligo-capping; ss. 06-NOV-2001 (first entry) CDNA; 1687 SEQ ID NO: 3459

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Matches 883
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           The invention relates to primers for synthesising full length cDNA clones. 830 cDNA molecules encoding a human protein have been isolated and nucleotide sequences of 5'- and 3'-ends of the cDNA molecules have been determined. Primers for synthesising the full length cDNA are useful for clarifying the function of the protein encoded by the cDNA. The full length clones were obtained by construction of full length enriched cDNA libraries that were synthesised by the oligo-capping method. The primers enable the production of the full length cDNA easily without any special methods. The present sequence is a full length human cDNA of the invention. Note: The sequence data for this patent did not form part of the printed specification, but was obtained in CD-ROM format directly
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Sequence
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genetic manipulation.
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su A, Sugiyama
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TTACTGACTGAAAACTTTTTTAAGAGATACTTGGTGTAATTTCCAGTTCTATACGTCCCTA
                     TTACTGACTGAAAACTTTTTAAGAGATACTTGGTGTAATTTCCAGTTCTATACGTCCCTA
                                                                                                GGCAGACAGCATTTACAGAATTTAGATGATGCTGTAAATGGGTCTGCATGGACAATCTTA
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Nagai K, Kojima
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Pred. No. 7e-207;
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S, Otsuki T, Koga
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This invention relates to a novel primers useful for synthesising full length cDNA molecules that encode human proteins. Specifically, it refers to secretory or membrane proteins that are potential therapeutic agents/ target molecules in the field of medicine, and in particular genes encoding proteins that are associated with signal transduction, glycoproteins and transcription. The present invention describes a method for efficiently cloning a full length human cDNA from both the 5' and 3' ends using the oligo-capping method. This polynucleotide sequence is a

S 밁 S 밁 S 문 δ 밁 á ₽ Ş

Wakamatsu

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Sugiyama

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Nishikawa

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Isogai T, na T, Nagai

Hayashi K, K, Kojima

Ishii S, S, Otsuki

Kawai T, K

Koga Ή,

oligonucleotide

primers

(830 cDNAs) useful

for

synthesizing

human

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3459;

1340pp; English.

2004-204755/20. DB; ADL31427.

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GAAAGCAGTTTCTAGGAAATGTTTAAATAAAAGAGAGTCTTCACCTT
                         GAAAGCAGTTTCTAGGAAATGTTTAAATAAAAGAGAGTCTTCACCTT
                                                                              GAAACATATAACATGTGGCTGGCTCTTGTTTTGTAAACCAAATGATTAATCTTCACTTGA
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07-JUL-2000;
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oligo-capping method; ss; gene.
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Best Local Similarity 93.4%;
Matches 498; Conservative
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11-JAN-2000;
02-MAY-2000;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     The invention relates to primers for synthesising full length cDNA clones. 830 cDNA molecules encoding a human protein have been isolated and nucleotide sequences of 5'- and 3'-ends of the cDNA molecules have been determined. Primers for synthesising the full length cDNA are useful for clarifying the function of the protein encoded by the cDNA. The full length clones were obtained by construction of full length enriched cDNA libraries that were synthesised by the oligo-capping method. The primers enable the production of the full length cDNA easily without any special methods. The present sequence is the nucleotide sequence of the 3'-end of a cDNA provided in the invention. Note: The sequence data for this patent did not form part of the printed specification, but was obtained in CD-ROM format directly from EPO
                                                                                                                                                                                                                                                                                                                                                                                                                                                                        Sequence 533 BP; 150 A; 103 C; 89 G; 175 T; 0 U; 16 Other;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Claim 3; SEQ ID NO 1475; 1380pp +
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  830 Primers useful for synthesizing
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su A, Sugiyama T, Nagai
                                                                        CTGAACAATCCCCCTTCCCCGAGAAAGGACTCCCCTTTGCCCTCCAAACCATCAATGCCTTA
                                                                                                                                                                                                                                                                                  CCATGTGGCAGACAGCATTTACAGAATTTAGATGATGCTGTAAATGGGTCTGCATGGACA
                                                                                                                                                                                                                                                                                                                               GTNCAGGATTTGNAANAAGNGGAANTTGGTATCAACCCGGGATTATTTTGNTTGAGATG
                                                                                                                        TCCCTAATGAACTCCGTTAACAGGCAGCATAAATACAATTNTGTTATACCCATGCGGCCC
                                                                                                                                          TCCCTAATGAACTCCGTTAACAGGCAGCATAAATACAACTCTGTTATACCCCATGCGGCCC
                                                                                                                                                                                           ATTTTATTACTGACTGAAAACTTTTTAAGANANACTNGGTGTAATTTCCAGTTNNATACG
                                                                                                                                                                                                                   ATCTTATTACTGACTGAAAACTTTTTTAAGAGATACTTGGTGTAATTTTCCAGTTCTATACG
                                                                                                                                                                                                                                                            CCATGTGGCAGACAGCATTTACAGAATNTAGATGATGCTGTAAAT-GGTCTGCATGGACA
                                                                                                                                                                                                                                                                                                                                                                     GTCCAGAATCTGCTACAAGATGACTTTGGTATCAAACCCGGAATAATCTTTGCTGAGATG 455
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             ; 99JP-00194486.
; 2000JP-00118774.
; 2000JP-00183765.
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                                                                                                                                                                                                                                                                                                                                                                                                    Score 476.2; DB 4;
Pred. No. 3.4e-107;
0; Mismatches 34;
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K, Kojima S,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Sequence Listing; English
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S, Otsuki T, Koga
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Length 533; Indels

1;

Gaps

414 515 473

695

294 635 354 575

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RESULT 7
ADL29442/c
ID ADL294
XX ADL294
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XX 20-MAY
XX Oligo-
XX Homo 8
PN EP1396
XX EP1396
XX EP1396
XX 08-JUL
XX 01-AT
PI Wakama
XX WPI; 2
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XX WPI; 2
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Matches 498
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11-JAN-2000; 2000JF-00118774.
02-MAY-2000; 2000JF-00183865.
                                                                                                                                                       This invention relates to a novel primers useful for synthesising full length cDNA molecules that encode human proteins. Specifically, it refers to secretory or membrane proteins that are potential therapeutic agents/ target molecules in the field of medicine, and in particular genes encoding proteins that are associated with signal transduction, glycoproteins and transcription. The present invention describes a method for efficiently cloning a full length human cDNA from both the 5' and 3' ends using the oligo-capping method. This polynucleotide sequence is the 3' end of a full length human cDNA sequence of the invention.
                                                                                                                            Sequence 533 BP;
                                                                                                                                                                                                                                                                                                                        Disclosure; SEQ ID NO 1475; 1340pp; English.
                                                                                                                                                                                                                                                                                                                                                       New oligonucleotide primers (830 cDNAs) useful length human cDNAs.
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                                                                                                                                                                                                                                                                                                                                                                                                                                         Wakamatsu
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oligo-capping method; ss.
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                                                                            Similarity
GTNCAGGATTTGNAANAAGNGGAANTTGGTATCAACCCGGGATTATTTTTGNTTGAGATG
                            GTCCAGAATCTGCTACAAGATGACTTTGGTATCAAACCCGGAATAATCTTTGCTGAGATG
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   ACTTGAGAAAGCAGTTTCTAGGAAATGTTTAAATAAAAGAGAGTCTTCACCTT
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                                                              Conservative
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a T, Nagai K, Kojima
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                                                             Score 476.2; D
Pred. No. 3.4e-
0; Mismatches
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                                                                              2; DB 12;
8.4e-107;
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T, Koga
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New polynucleotide and polypeptide useful for diagnosing, preventing or treating conditions such as neurodegenerative diseases, anemias, platelet

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RRESULT 8
ADC3230
ID ADC33
XX ADC32
XX ADC3
XX ADC3
XX Huma
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Human; diagnostic; drug screening; forensics; gene mapping; biodiversity assessment; Parkinson's disease; Alzheimer's disease; neurodegenerative diseases; anaemia; plattelet disorder; wound; burns; ulcers; osteoporosis; autoimmune disease; cancer; molecular weight marker; food supplement; antiparkinsonian; nootropic; neuroprotective; antianaemic; anticoagulant; thrombolytic; vulnerary; antiulcer; osteopathic; immunosuppressive; antiinflammatory; cytostatic; gene therapy; chromosome 5; ss.
                                                                                                                                                                 Tang
Zhou
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      Human novel cDNA contig sequence,
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                                                                    P-PSDB;
                                                                                                                                             Haley-Vicente
                                                                                                                                                                                                                                                                                      24-SEP-2001; 2001US-0324631P
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                                                                      2003-371981/35.
DB; ADC33147.
                                                                                                                                                            TY, Zhang J, P, Ghosh M,
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disorders, wounds, burns, ulcers, osteoporosis, autoimmune diseases or

SEQ ID NO 2462; 1185pp; English

The invention relates to 971 novel human cDNA sequences (ADC29919-CC ADC20889) and the polypeptides they encode (ADC30890-ADC31860). The CC invention also relates to nucleic acid sequences over 99% identical with CC the novel human cDNAs. The invention additionally encompasses expression CC vectors and host cells comprising a nucleic acid of the invention; the recombinant production of a polypeptide of the invention; an antibody against a polypeptide of the invention; an antibody CC against a polypeptide of the invention; an antibody CC identifying a compound which binds to a polypeptide of the invention. The CC invention further discloses methods of peventing, treating or ameliorating a medical condition; kits comprising polynuclectide probes CC and/or monoclonal antibodies for carrying out the methods of the contigs of the contigs equences corresponding to the cDNA sequences of the invention (ADC31861-ADC32627) and the polypeptides encoded by the contigs (ADC32628 - ADC3394). The nucleic acids and polypeptides of the invention are combined to the mutation of mutations responsible for genetic disorders or other craits, for assessing biodiversity, and in producing many other types of data and products dependent on DNA and amino acid sequences. They are CC disease and other neurodegenerative diseases, anaemia, platelet cancer. The nucleic acids may also be used as hybridisation probes or CC cancer. The nucleic acids may also be used as hybridisation probes or CC cancer. The nucleic acids may also be used as hybridisation probes or CC cancer. The nucleic acids may also be used as hybridisation probes or CC and as food supplements. The present sequence represents a human contig contig sequence used in an example of the invention. Note: The sequence data for this patent did not form part of the printed specification, but was CC this patent did not form part of the printed specification, but was ftp.wipo.int/pub/published_pct_sequences.

Sequence 522 BP; 157 A; 100 C; 129 G; 136 T; 0 U; 0 Other,

DB 10;

Length 522;

B á 밁 Ś 밁 á S 밁 밁 á 밁 S 밁 Query Match Best Local Similarity Matches 470; 430 352 370 292 310 232 250 172 190 112 52 70 AAATAAATTCCTGCCCTCTTTCTCTCTCTTTGGGGTAAAAGGCACAGTGTGGGATACAAGTC GAAATACATAAAGTTTTCCTCTTCTGCCTTGGATATTTATAATGGGTATCGGGAAGTCTA 111 GAAATACATAAAGTTTTTCCTCTTCTGCCTTGGATATTTATAATGGGTATCGGGAAGTCTA AACCCGGAATAATCTTTGCTGAGATGCCATGTGGCAGACAGCATTTACAGAATTTAGATG AAGATGACACAGATGAAGCCCTCAGAGTCCCAGAATCTGCTACAAGATGACTTTGGTATCA CAGGATATCATGAGTCAGATTCCAAGAAGTCTGAAGATCTATCCTTGTGTAATGTTGCTG CAGGATATCATGAGTCAGATTCCAAGAAGTCTGAAGATCTATCCTTGTGTAATGTTGCTG **AAGATGACACAGATGAAGCCCTCAGAGTCCAGAATCTGCTACAAGATGACTTTGGTATCA** <u>AACCCGGAATAATCTTTGCTGAGATGCCATGTGGCAGACAGCATTTACAGAATTTAGATG</u> Conservative 17.3%; 99.8%; ٥, Score 469.4; DB 10 Pred. No. 1.6e-105; 0; Mismatches 1; Indels ٥, 429 471 489 411 351 369 291 309 231 249 189 129 171

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RESULT 9
ABN96310/c
ID ABN963
XX ABN963
XX ABN963
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XX Gene;
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XX Metast
XX Metast
XX Metast
XX Homo E
XX WO2002
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XX Homo E
XX WO2002
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XX Homo E
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XX WO2002
XX CENT
PM 02-OCT
XX WPI;
PM (GENEXX WPI;
XX WPI;
XX WPI;
XX Claim
XX Claim Gene; liver cancer; ds; hepatocellular carcinoma; hepatotropic; metastatic liver tumour; cytostatic; expression profile; disease state; disease progression; drug toxicity; drug efficacy; drug metabolism. Horne D, WO200229103-A2 Gene #2808 used to diagnose liver cancer. 02-OCT-2000; 2000US-0237054P 02-OCT-2001; 2001WO-US030589 13-AUG-2002 (first entry ABN96310; ABN96310 standard; Homo sapiens (GENE-) GENE LOGIC INC Alvares C, DNA; Peres-Da-Silva S, Vockley JG; ВP

Diagnosing and detecting the progression of liver cancer, hepatocellular carcinoma or metastatic liver tumor in a patient, involves detecting the level of expression of two or more genes in a liver tissue sample.

Claim 1; SEQ ID NO 2808; 298pp; English.

The invention relates to a novel method for diagnosing and detecting the progression of liver cancer, hepatocellular carcinoma or metastatic liver tumour in a patient, and differentiating metastatic liver cancer from the patocellular carcinoma in a patient, involving detecting the level of the patocellular carcinoma in a patient, involving detecting the level of expression of two or more genes represented in ABN93503-ABN97455 in a cissue sample. The method of the invention has hepatotropic, and crissue static activity. The method is useful for diagnosing and detecting the progression of liver cancer, hepatocellular carcinoma and metastatic cliver carcinoma in a patient. The method is useful for identifying expression profiles which serve as useful diagnostic markers as well as markers that can be used to monitor disease states, disease progression, of drug toxicity, drug efficacy and drug metabolism. Note: The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from WIPO at fig.wipo.int/pub/published_pct_sequences Sequence 253 BP; 100 A; 36 C; 44 G; 70 T; 0 U; 3 Other

S 밁 ঠ 밁 Query Match
Best Local Similarity
Matches 243; Conserv 186 GATAGTAGCTTACTCACTCCTCTTTTTCAGTTTTCATAATAAGTATTCATTTTTTTG CCATAATGCTTCCTGTAAAGCCAATTTTATATATACTAATAAAACATGAACTGCCCACTCTT GATAGTAGCTTACTACTCACTTCTCTTTTTCAGTTTTTCATAATAAGTATTCATTTTTTTG CTTATTAGAGGAGAGAGGTTTTCATTGCATAGTGACATTTTCTTAGCCTTAACGTTCT CCATAATGCTTCCTGTAAAGCCCAATTTTATATACTAATAAAAACATGAACTGCCCACTCTT CNTATTAGAGGAGGAGATGGTTTTCATTGCANAGTGACATTTTCTTAGCCTTAACGNTCT Conservative 98.98; Score 243; DB Pred. No. 1e-4 0; Mismatches 0 1e-49; DB 6; Length 253; Indels <u>,</u> Gaps 127 2651 2591 187 67 0

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nucleotide sequence from a genome of an organism corresponds to a nucleotide sequence of an open reading frame; for preparing a contig, nucleic acid molecule from a genome of an organism; and for sequencing all or part of a genome of an organism. mRNA is obtained from mammalian or human cell which is associated with a pathological condition e.g. a colon cancer or breast cancer cell. The method is useful for analyses of populations of subjects and can be used to carry out genetic analyses of large or small populations. further, it can be used to study living systems to determine if e.g. there have been genetic shifts which render an individual or population more or less likely to be afflicted with diseases such as cancer, to determine antibiotic resistance or nontolerance, and so forth. The method can also be used in the study of congenital diseases, and the risk of affliction to a foctus, as well as the study of whether the conditions are likely to be passed to offspring
                                                                                                                                                                                                                                                        The invention describes a method of determining open reading frames in the genome of organism, comprising contacting mRNA from cell of organism with a single oligonuclectide primer (I) at Low stringency, preparing single-stranded cDNA by reverse transcribing mRNA with (I), amplifying cDNA, sequencing the product, and repeating the contacting, preparing and amplifying steps with different primers and sequencing resulting nucleic acids. The method is useful for: determining that a known nucleic acids.
                                                                                                                                                                                                                                                                                                                                                                                                                                                              Determining open reading frames of genome of an organism e.g. a human suffering from cancer involves use of single oligonucleotide primer a low stringency for preparing single-stranded cDNA from mRNA of
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Simpson AJG,
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                                                                                                                                                                                                                                                                                                                                                                                                          Example 9; Page 430; 959pp; English
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(NETO/)
(BREN/)
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Matches 270;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 through ova or sperm. The analyses for pathological conditions can be carried out in all animals, plants, birds, fish, etc. Using this method, in the area of agriculture, for example the genomes of food crops can be studied to determine if resistance genes are present, defects in plant genomes can also be studied in this way. Similarly, the method permits determination of the pathogens which integrate into the genome, such as retroviruses and other integrating viruses such as influenza virus, have undergone shifts or mutations, which may require different approaches to therapy. This method is also applied to eukaryotic method essentially eliminates sequencing of non-coding portions. This sequence represents a polynucleotide isolated from human colon cancer cell cDNA library
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Human biallelic polymorphic DNA fragment WI-11806
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Match 5.8%;
Local Similarity 73.6%;
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Pred. No. 3.7e-28;
D; Mismatches 82
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    660;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               15;
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RESULT 11

AAX10464/c

ID AAX10464 standard; DNA; 104 BE

XX

AC AAX10464;

XX

TO AAX10464;

XX

Polymorphism; biallelic; humar

XX

Polymorphism; biallelic typing;

XX

Metection; phenotypic typing;

XX

Mutoimmune disease; cancer; if

XX

OS Homo sapiens.

XX

PN

WO9820165-A2.

XX

PD

14-WAY-1998.

XX

PR

05-NOV-1997; 97WO-US020313.

XX

RR

06-NOV-1996; 96US-0030455P. Polymorphism; biallelic; human; forensic; paternity testing; disease; detection; phenotypic typing; characteristic; infection; hereditary; autoimmune disease; cancer; inflammation; drug; therapy; medicament; treatment; marker; ss. 97WO-US020313

96US-0030455P

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RESULT 12
ABN46564/c
ID ABN465
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         AXX10269-X12937 are human DNA fragments which contain biallelic CC polymorphic markers which have been isolated using the primers CC represented in AAX09121-X10268. The base occupying the polymorphic site is indicated by the appropriate IUPAC-IUB ambiguity code. These fragments CC can be used in methods for determining polymorphic forms in an individual C for use in e.g. forensics, paternity testing or for phenotypic typing for CC diseases such as agammaglobulinemia, diabetes insipidus, Jesch-Nyhan CC syndrome, muscular dystrophy, Wiskott-Aldrich syndrome, Fabry's disease, CC familial hypercholesterolemia, polycystic kidney disease, hereditary CC spherocytosis, von Willebrand's disease, tuberous sclerosis, hereditary comparting tetlangiectasia, familial colonic polyposis, Ehlers-Danios System, infection by pathogenic microorganisms, and characteristics such autoimmune diseases, inflammation, cancer, diseases of the nervous System, infection by pathogenic microorganisms, and characteristics such alongevity, appearance (e.g. baldness, obesity), strength, speed, CC andurance, fertility, and susceptibility or receptivity to particular CC drugs or therapeutic treatments. The isolated polymorphic nucleic acid CC segments can also be used to produce medicaments for the treatment or CC prophylaxis of such diseases
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Query Match
Best Local Sim
Matches 103;
                                                                                                                                                                                                                                                                                                 Human spliced transcript detection oligonucleotide SEQ ID NO:19312.
                                                                                                                                                                                                                                                                                                                                                                                                                                    ABN46564 standard; DNA;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              New isolated nucleic acid segments from the human genome - used fo determining polymorphic forms for use in e.g. forensics, paternity testing or phenotypic typing for disease.
28-JUL-2000; 2000US-0221607P.
02-MAY-2001; 2001US-0287724P.
                                                                                                                                                    WO200210449-A2
                                                                                                                                                                                            Homo sapiens
                                                                                                                                                                                                                                                                                                                                              15-JUL-2002
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       Sequence 104 BP; 34 A; 15 C;
                                                              20-JUL-2001; 2001WO-18001903
                                                                                                                                                                                                                                       splice
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      Claim 1; Page 53; 310pp; English.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           (WHED ) WHITEHEAD INST BIOMEDICAL
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               2652 CATGCCTGCCAAACTTGGGGCAATTGATGCTAAATGGTATTTTT 2695
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         2592 CCATAATGCTTCCTGTAAAGCCAATTTTATATACTAATAAAACATGAACTGCCCACTCTT
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                                                                                                                                                                                                                                       mouse; rat; splice transcript; detection; RNA transcript variant; transcriptome; oligonucleotide library; ss.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        Similarity
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         CATGCCTGCCAAACTTGGGGCAATTGATGCTAAATGGTATTTTT
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              CATAATGCTTCCTGTAAAGCCAATTTTATATACTAATAAACMTGAACTGCCCACTCTT
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                                                                                                                                                                                                                                                                                                                                              (first entry)
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Pred. No. 1.9
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      Mismatches
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       31 T;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       U; 1 Other;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      Indels
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The present invention describes oligonuclectide libraries for detecting CC messenger RNAs that populate a (sub-)transcriptome, where the (sub-)transcriptom comprises messenger RNAs transcribed from multiple CC transcription units that populate a genome. The library comprises several CC oligonuclectides, each capable of hybridising selectively to a set of CC messenger RNAs transcribed from a given transcription unit of the genome, which encodes one or more messenger RNA splice variants. The CC oligonuclectide libraries are useful for detecting mRNAs from a CC quantitatively characterising the corresponding transcriptome, and in CC quantitatively characterising the corresponding transcriptome, and in CC transcriptomes. The libraries may also be used as specialised mini CC transcriptomes. The libraries may also be used as specialised mini CC transcriptomes. The libraries of a sub-transcriptome under a particular biological or pathological state, and so allowing the detection of tissue c and pathology-specific genes such as those genes only expressed in CC specific tissue under a specific pathological condition; to detect evariants of a transcriptome of a patient suffering from a particular CC developmental specific genes; and to detect RNA transcripts and splice variants of a transcriptome of a patient suffering from a particular cate, humans and mice, which are used in the exemplification of the grate of the printed specification, but was obtained in electronic format condition; from wifo at ftp.wipo.int/pub/published_pct_sequences
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        New oligonucleotide libraries comprising oligonucleotides which selectively hybridize to mRNAs transcribed from a transcription unit of genome, useful for detecting tissue-, pathology-, and developmental-
Sequence 60 BP; 24 A; 10 C; 12 G; 14 T; 0 U; 0 Other;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         Example 1; SEQ ID NO 19312; 47pp; English
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                specific genes.
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ABL32333;
                                    ABL32333 standard;
                                    DNA; 8895
                                    BP
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Human

immune

system associated gene SEQ

ID NO:

306.

26-MAR-2002

(first entry)

맑 á

6

Matches Query Match Best Local (

59;

Conservative

0

Similarity

2.1%;

Score 58.4; DB 6; Pred. No. 0.00023; 0; Mismatches 1

DB 6;

Length Indels

0

Gaps

0

2490 GGTTTTCATTGCATAGTGACATTTTCTTAGCCTTAACGTTCTGATAGTAGTAGCTTACTACTC

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antiarteriosclerotic; antianaemic; cytostatic; nootropic; neuroprotective; anti-HIV; anticonvulsant; ontihalmological; antirheumatic; antiarthritic; antidiabetic; antiastic; antiarthritic; antidiabetic; antieriosclerosis; anaemia; antiinflammatory; cancer; eye disease; arteriosclerosis; anaemia; acute myeloid leukaemia; Alzheimer's disease; AIDS; epilepsy; neurofibromatosis; rheumatoid arthritis; psoriasis; bowel disease
                                                                                                                                                                                                                                                                                                                                                                                                                               Human; immune system disease; cytosine methylation; antiasthmatic;
Homo sapiens.
                                                                                                                                             epilepsy;
bowel disease;
                                                                                                                                                  gene;
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02-JUL-2001; 2001WO-EP007537

03-JAN-2002.

WO200200928-A2

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RESULT 14
ABL33879
ID ABL33
XX
AC ABL33
XX
DT 26-MP
XX
DB Humar
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Query Match
Best Local Similarity
Matches 203; Conserv
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Sequence 8895 BP; 2269 A; 62 C; 1898 G; 4665 T; 0 U; 1 Other;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          30-JUN-2000; 2000DE-01032529.
01-SEP-2000; 2000DE-01043826.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      macular degeneration, arteriosclerosis, anaemia, cancer, acute mys leukaemia, Alzheimer's disease, AIDS, epileppy, neurofibromatosis, rheumatoid arthritis, psoriasis and inflammatory/ulcerative bowel diseases. The present sequence is a gene of the invention
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       The present invention provides a number of human immune system associated genes which are modified by the methylation of cytosines. The sequences can be used in the diagnosis and treatment of immune system disorders, including eye diseases such as retinopathy, neovascular glaucoma and
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       Claim 1; SEQ ID NO 306; 32pp +
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Nucleic acid comprising diagnosis and treatment
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  WPI; 2002-130909/17.
                                       26-MAR-2002
                                                                                       ABL33879 standard; DNA; 5378
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                                                                                                                                                                                                                                                                                                        ATGCTGTCATTACACTTAMAGTCACAGGAAAAAAATATACATGCTTACTCAGGCTTTCTT
                                                                                                                                                    TTTTGTTTTTT 8745
                                                                                                                                                                             TTCATTTTTT 2590
                                                                                                                                                                                                      CCTTAACGTTCTGATAGTAGCTTACTACTCACTTCTCTTTTTCAGTTTTCATAATAAGTA
                                                                                                                                                                                                                                                                                                                                ATTCCCCACTTGTATATCCCCTACCAGTACCGGGATCTGCACACATCTTTTTGCAGTTAC
                                                                                                                                                                                                                                                                                                                                                                                                             AAAAATAAATTTTTATAGAGATCCTTGAGTAAAGACATTTTGCTTAATTTCTTTTTCTT
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              system
                                       (first
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        2.1%;
47.1%;
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                                       entry)
              associated
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            fragment of chemically modified gene, of diseases associated with abnormal
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             gene
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       Sequence Listing;
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               SEQ
               IJ
               ö
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       Length 8895;
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                                                                                       RESULT 15
ABK31249
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Best Local Sim
Matches 118;
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   Signal transduction associated
                           23-APR-2002
                                                                            ABK31249
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01-SEP-2000;
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DNA;

10543

gene modified complementary DNA #46.

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The present invention provides a number of human immune system associated genes which are modified by the methylation of cytosines. The sequences can be used in the diagnosis and treatment of immune system disorders, including eye diseases such as retinopathy, neovascular glaucoma and macular degeneration, arteriosclerosis, anaemia, cancer, acute myeloid leukaemia, Alzheimer's disease, AIDS, epilepsy, neurofibromatosis, rheumatoid arthritis, psoriasis and inflammatory/ulcerative bowel diseases. The present sequence is a gene of the invention
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Nucleic acid comprising diagnosis and treatment methylation.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Human; immune system disease; cytosine methylation; antiasthmatic; antiarteriosclerotic; antianaemic; cytostatic; nootropic; neuroprotective; anti-HIV; anticonvulsant; ophthalmological; antirheumatic; antiarthritic; antidiabetic; antisporiatic; antiarthritic; antidiabetic; antisporiatic; anaemia; anteriosclerosis; anaemia; acute myeloid leukaemia; Alzheimer's disease; AIDS; epilepsy; neurofibromatosis; rheumatoid arthritis; psoriasis; bowel disease;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Sequence 5378 BP; 1543 A; 134 C; 1135 G; 2566 T; 0 U; 0 Other;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    02-JUL-2001; 2001WO-EP007537
                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Local Similarity
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                                                                                                                                                                                                                                                                                                                                                                   ARARARGTTTGGATTCGATRARAGTGATATGTARTAGTTGCAGARGGCTTTATATATGCTT
                                                                                                                                                  ACTTAAAGTCACAGGAAAAAATATACATGCTTACTCAGGCTTTCTTAAAAAATATTTT
                                                                                                                                                                                                                   TATAGAGATCCTTGAGTAAAGACATTTTGCTTAATTTCTTTTTTCTTATT
                                                                                                           1.9%;
ilarity 51.3%;
Conservative
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2000DE-01043826.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                             Score 50.8;
Pred. No. 0.
                                                                                                                                                                                                                                                                                                                                                                                                                                    Mismatches
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         DB 6;
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                                                                                                                                                                                                                                                                                                                                                                                                                                               The present invention relates to chemically modified DNA sequences of Signal transduction associated genes. The DNA sequences are chemically comodified using a solution of bisulphite, hydrogen sulphite or disulphite. Also disclosed are oligonucleotides and/or pNA oligoners for detecting the cytosine methylation state (CpG islands) of these genes, and a method cf or the diagnosis and/or therapy of genetic and epigenetic parameters of genes associated with signal transduction. The genomic DNA, can be compared to the contain DNA, e.g. cell clines, bloppies, blood, sputum, stool, urine, cerebral-spinal fluid, ctisque embedded in paraffin such as tissue from eyes, intestine, kidney, can be cuseful for the diagnosis and therapy of diseases associated with signal cuseful for the diagnosis and therapy of diseases associated with signal chemically pretreated genomic DNA sequences of the invention are chemically pretreated genomic DNA sequences of different genes associated with signal transduction, or their complementary sequences. Note: The sequence data for this patent did not form part of the printed contains. The securities associated with signal transduction, or their complementary sequences. Note: The sequence data for this patent did not form part of the printed contains the printed specification, but was obtained in electronic format directly from the contact.
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Best Local Simi
Matches 221;
                                                                                                                                                                                                                                                                                                                                                                                                                    Sequence 10543 BP; 2870 A; 177 C; 2279 G; 5217 T; 0 U; 0 Other;
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                                                                               AATAAATTTTTATAGAGATCCTTGAGTAAAGACATTTTGCTTAATTTCTTTTTTCTTATT
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740 TTTRNMTTSTNMTNNNNNMWACTNNNNNMWKAYYAHATNNWGCWWNNTDARRTNNTTV 681

1245 GGATGTTGCAATAAGCATTCGGGTACTATCACCCAGAATATGAATTGCCAGAATAGAAC 1304

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800 HYT	1005 AG1	945 CAC ::: 860 YHF	latch cal Simi 81;	B 4	SULT 1 Sequence 22, Application US/09 Patent No. 6794342 GENERAL INFORMATION: APPLICANT: The University of TITLE OF INVENTION: Regulation FILE REPERENCE: 4810-58741 CURRENT APPLICATION NUMBER: U CURRENT FILING DATE: 2001-04 PRIOR APPLICATION NUMBER: US PRIOR APPLICATION NUMBER: US PRIOR APPLICATION NUMBER: US PRIOR APPLICATION STEEL US PRIOR APPLICATION STEEL US PRIOR APPLICATION ON STEEL US PRIOR APPLICATION STEEL US PRIOR APPLIC		41.8 41.8 41.6		42.2 42.2						43
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AAARMARTO TCAATCO	CAGAGTTT	PCTGCAGGI NNNNNNNNN	Scc Pre	ce B Bequence	:-708B-22/c :e 22, Application US/09806708B NO. 6784342 INFORMATION: ANT: The University of British Colum OF INVENTION: Regulation of Embryoni EFERENCE: 4810-58741 TAPPLICATION NUMBER: US/09/806,708B TF FILING DATE: 2001-04-03 APPLICATION NUMBER: US 60/147,133 FILING DATE: 1999-08-04 OF SEQ ID NOS: 23 RE: Patentin version 3.0 NO 22	ALI	US-09- US-09- US-09-	US-09-	US-09-	US-09-	US-07-	US-09-	-60-Sn	US-09-	-60-Sn
HYTTÄNABBCYRÄNNNNAÄARMARTCNNYMHAAVTTTHTDWCYKTWMVTWYWDMYTTMBT GATATGAAAAAGGAATTGTAATTGTATTGTTTTTTTTT	AGTAAACTTGACATTGTCAAGGAGTTTCAAGGACTTTTCCTTTCACAATTTTTCCTAGGTTCATG	CACAAGAAAGATAAATTTCTGCAGGACAGCCTATAAAATTGTGGGTACTTTTTGATGTTTC ::: :::: ::: !: YHHAARRWMNAWWTRTNNNNNNNNNNNNACRNTRTWWABWKHSWCNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNN	50.8; DB 4 No. 0.0012; matches 39	of A.t.,	onic	ALI GNMENTS	US-09-949-016-1237 US-09-806-708B-23 US-09-790-988-1 US-09-129-112-3	US-09-949-016-13294 US-09-949-016-13845	917-265A-104 917-265A-106	949-016-1183 949-016-1237	715-7518-2 949-016-1501	949-016-1516	US-09-949-016-1200 US-09-949-016-1705	806-708B-23	949-016-1600
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Patent No. 6784342

GENERAL INFORMATION:
APPLICANT: The University of British Columbia
TITLE OF INVENTION: Regulation of Embryonic Transcription in Plants
FILLE REFERENCE: 4810-58741
CURRENT APPLICATION NUMBER: US/09/806,708B
CURRENT APPLICATION NUMBER: US/09/806,708B
CURRENT FILING DATE: 2001-04-03
PRIOR APPLICATION NUMBER: US 60/147,133
PRIOR FILING DATE: 1999-08-04
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                                                                                                                                                                                                                                                                                                                  Query Match
Best Local :
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SOFTWARE: PatentIn version 3.0
                                                                                                                                                                                                                                                                                                                                                                                          NAME/KEY: promoter LOCATION: (1)..(114 OTHER INFORMATION:
                                                                                                                                                                                                                                                                                                                                                                                                                                                      TYPE: DNA ORGANISM: Artificial sequence FEATURE:
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                    TCTGTGAAATGAAAGGGTTGAATTGATGGATCTCTAAAGGCTTTTGTCCTCTATGAGGAT
                                                                                                                                     GWRHRYWRWRAMBDTVDHHYYTAMNNAWTTWCMMDKDDKRTRWWWKKNNNATGWDDDTKY
                                                         HMWNNNGCBTVTWMVRYKTDRDWSBKRMNYGMBWWKNWSYDVTYYWWVWDDMCKRKVRRW
                                                                                                AGTCCAATTTCTGCCACTGAGGATGAATGTAACTGTGGGCAAACTATTTACCCTCCTTTA
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                                                                                                                                                                                                                                                                                            3%; Score 50.6; DB 4;
k; Pred. No. 0.0014;
306; Mismatches 412;
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Patent No. 5670367
                                                                                                                                                                                                                                                                                                                                                                                                             GENERAL INFORMATION:
APPLICANT: DORNER, F.
APPLICANT: SCHEIFLINGER, F.
APPLICANT: FALKNER, F. G.
APPLICANT: FALKNER, F. G.
TITLE OF INVENTION: RECOMBINANT FOWLPOX VIRUS
                                                                                              ZIP: 22313-0299
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC Compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PATENTIN Release #1.0, V
CURRENT APPLICATION DATA:
APPLICATION UMBER: US/08/232,463
FILING DATE:
FILING DATE:
                                         PRIOR APPLICATION DATA APPLICATION NUMBER:
                                                                                                                                                                                                                                                                                                                                                                          CORRESPONDENCE ADDRESS:
                                                                                                                                                                                                                                                                                                                                                                                                   NUMBER OF SEQUENCES:
      FILING DATE:
APPLICATION NUMBER:
                                                                                                                                                                                                                                                                       COUNTRY:
                                                                                                                                                                                                                                                                                                  STATE:
                                                                                    CLASSIFICATION:
                                                                                                                                                                                                                                                                                                                    CITY: Alexandria
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SEQUENCE CHARACTERISTICS:
LENGTH: 7218 bann
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Best Local Similarity
Matches 9; Conserv
                                                                                                                                                                                                                                    GENERAL INFORMATION:
ZIP: 22313-0299
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
                                                                                                                                                           APPLICANT: DORNER, F.
APPLICANT: SCHEIFLINGER, F.
APPLICANT: FALKNER, F. G.
TITLE OF INVENTION: RECOMBINANT
NUMBER OF SEQUENCES: 52
                                                                                                               CORRESPONDENCE ADDRESS:
ADDRESSEE: Foley & Lardner
STREET: 1800 Diagonal Road
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REGISTRATION NUMBER: 29,768
REFERENCE/DOCKET NUMBER: 30
TELECOMMUNICATION INFORMATION:
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ATTORNEY/AGENT INFORMATION:
                                                                        COUNTRY:
                                                                                                   CITY: Alexandria
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TYPE: nucleic acid
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TELEPAX: (703)683-4109
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2.5%; Pred. No. 0.0069;
ative 213; Mismatches 145; Indels 0
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US-08-232-463-14
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INFORMATION FOR SEQ ID NO:
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 ATTORNEY/AGENT INFORMATION:
NAME: BENT, Stephen A.
REGISTRATION NUMBER: 29,768
REFERENCE/DOCKET NUMBER: 3047
TELECOMMUNICATION INFORMATION:
TELEPHONE: (703)836-9300
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          IMMEDIATE SOURCE:
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              SEQUENCE CHARACTERISTICS:
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CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/232,463
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      'Match 1.8%; Score 48.6; DB 1; Local Similarity 0.8%; Pred. No. 0.015; les 3; Conservative 216; Mismatches 140;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     TELEFAX:
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REFERENCE TO THE PROPERTY OF T
                                                                     GTGGCAGACAGCATTTACAGAATTTAGATGATGCTGTAAATGGGTCTGCATGGACAATC 518
                                                                                                                                                                                                                                AGAATCTGCTACAAGATGACTTTGGTATCAAACCCGGAATAATCTTTTGCTGAGATGCCAT 459
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RESULT 5
US-09-949-016-15854/c
US-09-949-016-15854/c
; Sequence 15854, Application US/09949016
; Patent No. 681239
; GENERAL INFORMATION:
GENERAL INFORMATION: Craig et al.
TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES
TITLE OF INVENTION UNMBER: US/09/949,016
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT EPILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/231,755
PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498

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345 AAATATATATATAATTTTTTTTTT 371

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PRIOR APPLICATION NUMBER: 60/241,755
PRIOR FILING DATE: 2000-10-20
PRIOR APPLICATION NUMBER: 60/237,768
PRIOR FILING DATE: 2000-10-03
PRIOR APPLICATION NUMBER: 60/231,498
PRIOR APPLICATION NUMBER: 60/231,498
PRIOR FILING DATE: 2000-09-08
PRIOR FILING DATE: 2000-09-08
PRIOR PRIOR DATE: 207012
SOFTWARE: FASESEQ for Windows Version 4.0
SEQ ID NO 54021
PRIOR FILENCE
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     ; LENGTH: 55195
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-15854
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US-09-949-016-54021
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NUMBER OF SEQ ID NOS: 207012
SOFTWARE: FastSEQ for Windows Version 4.0
SEQ ID NO 15854
                                                                                                                                                                                                                                                                                                             Query Match
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ORGANISM: Human
-09-949-016-54021
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TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
FILE REFERENCE: CL001307
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CURRENT FILING DATE: 2000-04-14
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Local Similarity 53.5%;
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2310 AAAGACATTTTGCTTAATTTCTTTTTT 2336
                                                                                                                                                                                                                                 2130 ATAMAGTGATATGTAATAGTTGCAGAAGGCTTTATATATGCTTATAATGAAAAGATATTT
                                                                                                                al Similarity
107; Conserv
                                                                           AAAAATATACATGCTTACTCAGGCTTTCTTAAAAATTAAATTTTTATAGAGATCCTTGAGT 2309
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                                       TTTGTATATTGACAGCATAATTTATTTTAATGCTGTCATTACACTTAAAGTCACAGGAA
                                                                                                                                                                                          1.7%;
llarity 51.7%;
Conservative
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                                                                                                                                                                                                                                                                        Score 47; DB
Pred. No. 0.00
0; Mismatches
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Pred. No. 0.085;
0; Mismatches 87;
                                                                                                                                                                                                                                                                                           DB 4; Length 601; 0.0095;
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APPLICATION NUMBER: 60/237,768
PRIOR APPLICATION NUMBER: 60/231,498
PRIOR APPLICATION NUMBER: 60/231,498
PRIOR FILING DATE: 2000-09-08
PRIOR FILING DATE: 20
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FILE REPERENCES: CLOOI307

CURRENT APPLICATION NUMBER: US/09/949,016

CURRENT FILING DATE: 2000-04-14

PRIOR APPLICATION NUMBER: 60/241,755

PRIOR FILING DATE: 2000-10-20

PRIOR APPLICATION NUMBER: 60/237,768

PRIOR FILING DATE: 2000-10-03

PRIOR FILING DATE: 2000-10-03

PRIOR FILING DATE: 2000-09-08

NUMBER OF SEQ ID NOS: 207012

SOFTWARE: FASTSEQ for Windows Version 4.0

SEQ ID NO 54022

LUGGTH: 601
LENGTH: 601
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-54024
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US-09-949-016-54022
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US-09-949-016-54022
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APPLICANT: VENTER, J. Craig et al.
TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        Patent No. 6812339
GENERAL INFORMATION:
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Query Match
Best Local Similarity
Matches 107; Conserv
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Sequence 54024,
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RESULT 10
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US-09-949-016-108655
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                                                                                                                                                                                                                                                                                                                                      SEQ ID NO 108655
LENGTH: 601
TYPE: DNA
ORGANISM: Human
GENERAL INFORMATION:
           Sequence 108656, Application US/09949016 Patent No. 6812339
                                                                                                                                                                                                                                                                          Query Match 1.7
Best Local Similarity 51.7
Matches 107; Conservative
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             GENERAL INFORMATION:
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Best Local Similarity 51.7%;
Matches 107; Conservative
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                                                                                                                               AAATATATATATAATTTTTTTTTTT 371
                                                                                                                                                                              AAAGACATTTTGCTTAATTTCTTTTT 2336
                                                                                                                                                      AAAAATATACATGCTTACTCAGGCTTTCTTAAAAATTAAATTTTTTATAGAGATCCTTGAGT 2309
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Pred. No. 0.0095;
0; Mismatches 100; Indels
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OF DETECTION AND USES THEREOF
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GENERAL INFORMATION:

APPLICANT: VENTER, J. Craig et al.

TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASS;

TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF

FILE REFERENCE: CL001307

CURRENT APPLICATION NUMBER: US/09/949,016

CURRENT FILING DATE: 2000-04-14

PRIOR APPLICATION NUMBER: 60/241,755

PRIOR APPLICATION NUMBER: 60/237,768

PRIOR PILING DATE: 2000-10-03

PRIOR APPLICATION NUMBER: 60/231,498

PRIOR APPLICATION NUMBER: 60/231,498

PRIOR APPLICATION NUMBER: 60/231,498

PRIOR APPLICATION NUMBER: 60/231,498

PRIOR APPLICATION NUMBER: 00/231,498

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                                                                                                                 Query Match
Best Local Similarity
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LENGTH: 601
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TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
FILE REFERENCE: CLOO1307
CURRENT APPLICATION NUMBER: US/09/949,016
CURRENT FILING DATE: 2000-04-14
PRIOR APPLICATION NUMBER: 60/241,755
PRIOR APPLICATION NUMBER: 60/237,768
PRIOR APPLICATION NUMBER: 60/237,768
PRIOR APPLICATION NUMBER: 60/231,768
PRIOR FILING DATE: 2000-10-03
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107; Conserv
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ATAAAGTGATATGTAATAGTTGCAGAAGGCTTTATATATGCTTATAATGAAAAGATATTT 2189
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                                                                                      Conservative
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51.7%;
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                                                                              Score 47; DB 4; Length 601
Pred. No. 0.0095;
0; Mismatches 100; Indels
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OF DETECTION AND USES THEREOF
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US-09-949-016-14731, Application US/09949016; Sequence 14731, Application US/09949016; Patent No. 6812339; GENERAL INFORMATION: POLYMORPHISMS IN KNOWN GENES; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES; FILE REFERENCE: CL001307
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US-09-949-016-13303
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PRIOR APPLICATION NUMBER: 60/241,755
PRIOR FILING DATE: 2000-10-20
PRIOR APPLICATION NUMBER: 60/237,768
PRIOR APPLICATION NUMBER: 60/237,768
PRIOR FILING DATE: 2000-10-03
PRIOR PILING DATE: 2000-09-08
PRIOR FILING DATE: 2000-09-08
PRIOR FILING DATE: 2000-09-08
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APPLICANT: VENTER, J. Craig et al.
APPLICANT: VENTER, J. Craig et al.
TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
FILE REFERENCE: CL001307
CURRENT APPLICATION NUMBER: US/09/949,016
CURRENT FILING DATE: 2000-04-14
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Query Match
Best Local Similarity
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LENGTH: 104520
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FEATURE:
NAME/KEY: misc_feature
LOCATION: (1)...(104520)
OTHER INFORMATION: n = A,T,C
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                 ASSOCIATED
OF DETECTION AND
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                     THEREOF
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Sequence 54023, Application US/09949016

Fatent No. 6812339

GENERAL INFORMATION:

APPLICANT: VERITER, J. Craig et al.

TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED

TITLE OF INVENTION: WITH HUMAN DISBASE, METHODS OF DETECTION AND USES THEREOF

FILE REFERENCE: CLOO1307

CURRENT APPLICATION NUMBER: US/09/949,016

CURRENT FILING DATE: 2000-04-14

PRIOR APPLICATION NUMBER: 60/241,755

PRIOR APPLICATION NUMBER: 60/237,766

PRIOR FILING DATE: 2000-10-03

PRIOR FILING DATE: 2000-09-08

PRIOR APPLICATION NUMBER: 60/231,498

PRIOR APPLICATION NUMBER: 60/231,498

PRIOR FILING DATE: 2000-09-08
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CURRENT FILING DATE: 2000-04-14
PRIOR APPLICATION NUMBER: 60/241,755
PRIOR FILING DATE: 2000-10-20
PRIOR PILING DATE: 2000-10-20
PRIOR PILING DATE: 2000-10-03
PRIOR APPLICATION NUMBER: 60/237,768
PRIOR FILING DATE: 2000-09-08
PRIOR FILING DATE: 2000-09-08
PRIOR FILING DATE: 2000-09-08
NUMBER OF SEQ ID NOS: 207012
SOPTWARE: FRASESEQ for Windows Version 4.0
SEQ ID NO 14731
LENGTH: 126029
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                                                                                                                          ; ORGANISM: Human
US-09-949-016-54023
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; LOCATION: (1)...(126029)
; OTHER IMPORMATION: n = A,T,C
US-09-949-016-14731
                                                                                                                                                                                NUMBER OF SEQ ID NOS: 207012
SOFTWARE: FASTSEQ for Windows Version
SEQ ID NO 54023
LENGTH: 601
                                                               Query Match
Best Local Similarity
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Matches 107;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Query Match 1.7%;
Best Local Similarity 51.7%;
                                                                                                                                                                TYPE: DNA
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  AAATATATATATAATTTTTTTTTTT 42838
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                                            Score 46.6; DB 4;
Pred. No. 0.012;
1; Mismatches 100;
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Pred. No. 0.23;
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PRIOR APPLICATION NUMBER: 60/241,755
PRIOR FILING DATE: 2000-04-14
PRIOR PLICATION NUMBER: 60/241,755
PRIOR FILING DATE: 2000-10-20
PRIOR PILING DATE: 2000-10-03
PRIOR APPLICATION NUMBER: 60/237,768
PRIOR FILING DATE: 2000-09-08
PRIOR PILING DATE: 5007012
SOFTWARE: FastSEQ for Windows Version 4.0
SEQ ID NO 108657
LENGTH: 601
TYPE: DNA
ORGANISM: Human
US-09-949-016-108657
Search completed: September 23, 2005, 08:36:39 Job time: 449 secs
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US-09-949-016-108657
, Sequence 108657, Application US/09949016
, Patent No. 6812339
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Best Local Similarity 51.2%;
Matches 106; Conservative
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APPLICANT: VENTER, J. Craig et al.
TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
FILE REFERENCE: CL001307
CURRENT APPLICATION NUMBER: US/09/949,016
CURRENT FILING DATE: 2000-04-14
                                                                                                                                                                                                                                                                                                                                                     2130 ATAAAGTGATATGTAATAGTTGCAGAAGGCTTTATATATGCTTATAATGAAAAGATATTT 2189
                                                                                                           2310 AAAGACATTTTGCTTAATTTCTTTTTT 2336
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